

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L15	55957	antisense or triplex or ribozyme or RNAi or siRNA or "RNA interfer?" or "short interfer? RNA"	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 10:58
L16	186	"human x-linked inhibitor of apoptosis" or hiap-1 or xiap or "x-linked iap" or "miha protein"	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 10:59
L17	102723	cancer? or tumor? or carcinoma? or tumour?	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 11:00
L18	43	"follicular atresia?" or "atresia? follicular"	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 11:00
L19	8642	"inflammatory disorder"	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 12:04
L20	124	L15 AND L16	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 11:01
L21	96	L20 AND L17	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 11:02
L22	1	L20 AND L18	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 12:02
L23	5	L20 AND L19	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 12:03
L24	812	"inflamm? disease" or "inflamm?" or "inflamm? disorder"	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 12:04
L25	23	"inflammat? disease" or "inflammat?" or "inflammat? disorder"	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 12:04
L26	111793	inflammatory	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 12:04

L27	66	L20 AND L26	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2004/11/15 12:05
-----	----	-------------	---	----	----	------------------

Set	Items	Description
S1	4667	XIAP OR HIAP OR HILP OR MIHA
S2	213803	ANTISENSE OR RIBOZYME OR TRIPLEX OR SIRNA OR RNAI OR (SHORT INTERFER? RNA)
S3	7369699	CANCER OR TUMOR OR CARCINOMA
S4	1106	FOLLICULAR ATRESIA?
S5	1627772	INFLAMMAT?
S6	287	S1 AND S2
S7	89	RD S6 (unique items)
S8	61	S7 AND S3
S9	61	RD S8 (unique items)
S10	0	S7 AND S4
S11	3	S7 AND S5

? t s8/free/all

>>>"FREE" is not a valid format name in file(s): 399

8/6/1 (Item 1 from file: 5)

0015002823 BIOSIS NO.: 200400373612

Downregulation of Bcl-2, FLIP or IAPs (XIAP and survivin) by siRNAs sensitizes resistant melanoma cells to Apo2L/TRAIL-induced apoptosis
2004

8/6/2 (Item 2 from file: 5)

0014973429 BIOSIS NO.: 200400344218

Enhancement of C2-ceramide antitumor activity by small interfering RNA on X chromosome-linked inhibitor of apoptosis protein in resistant human glioma cells
2004

8/6/3 (Item 3 from file: 5)

0014959816 BIOSIS NO.: 200400330602

X-linked inhibitor of apoptosis protein inhibition induces apoptosis and enhances chemotherapy sensitivity in human prostate cancer cells
2004

8/6/4 (Item 4 from file: 5)

0014945078 BIOSIS NO.: 200400315835

Cyr61 expression confers resistance to apoptosis in breast cancer MCF-7 cells by a mechanism of NF-kappaB-dependent XIAP up-regulation
2004

8/6/5 (Item 5 from file: 5)

0014905560 BIOSIS NO.: 200400276317

Specific downregulation of bcl-2 and xIAP by RNAi enhances the effects of chemotherapeutic agents in MCF-7 human breast cancer cells
2004

8/6/6 (Item 6 from file: 5)

0014846985 BIOSIS NO.: 200400217040

Regulation and targeting of antiapoptotic XIAP in acute myeloid leukemia.
2003

8/6/7 (Item 7 from file: 5)
 0014818352 BIOSIS NO.: 200400186038
Akt phosphorylation and stabilization of X-linked inhibitor of apoptosis protein (XIAP).
 2004

8/6/8 (Item 8 from file: 5)
 0014795073 BIOSIS NO.: 200400162414
Mechanisms of apoptosis induction by a novel ring-substituted diindolylmethane (1,1-bis (3'-(5-methoxy indolyl))-1-(p-t-butylphenyl) methane) (BMITM) in leukemic cells.
 2003

8/6/9 (Item 9 from file: 5)
 0014780228 BIOSIS NO.: 200400146889
The triterpenoid CDDO-imidazolide induces apoptosis of CLL B-cells, through a Bcl-2-independent mechanism and synergizes with fludarabine.
 2003

8/6/10 (Item 10 from file: 5)
 0014730212 BIOSIS NO.: 200400100969
Induction of cIAP-2 in human colon cancer cells through PKCdelta/NF-kappaB.
 2003

8/6/11 (Item 11 from file: 5)
 0014727376 BIOSIS NO.: 200400098133
RNAi mediated downregulation of bcl-2 and xIAP may have therapeutical potential in human breast adenocarcinoma.
 2003

8/6/12 (Item 12 from file: 5)
 0014581909 BIOSIS NO.: 200300538099
Proteasome inhibitors potentiate leukemic cell apoptosis induced by the cyclin-dependent kinase inhibitor flavopiridol through a SAPK/JNK- and NF-kappaB-dependent process.
 2003

8/6/13 (Item 13 from file: 5)
 0014387492 BIOSIS NO.: 200300346211
The histone deacetylase inhibitor MS-275 promotes differentiation or apoptosis in human leukemia cells through a process regulated by generation of reactive oxygen species and induction of p21CIP1/WAF1.
 2003

8/6/14 (Item 14 from file: 5)
 0014380229 BIOSIS NO.: 200300336972
Bcl-2 Antisense Treatment for Waldenstrom's Macroglobulinemia.
 2002

8/6/15 (Item 15 from file: 5)
0014080092 BIOSIS NO.: 200300038811
Role of XIAP in the malignant phenotype of transitional cell cancer (TCC) and therapeutic activity of xiap antisense oligonucleotides against multidrug-resistant TCC in vitro.
2003

8/6/16 (Item 16 from file: 5)
0013916537 BIOSIS NO.: 200200510048
A novel tri-specific antisense oligonucleotide (AO) to the Inhibitors of Apoptosis Proteins (IAPs) enhances apoptosis in PC3 prostate cancer cells
2002

8/6/17 (Item 17 from file: 5)
0013815621 BIOSIS NO.: 200200409132
Antisense oligonucleotides targeting XIAP induce apoptosis and enhance therapeutic activity against human lung cancer cells when combined with anticancer drug in vitro and in vivo
2002

8/6/18 (Item 18 from file: 5)
0013365117 BIOSIS NO.: 200100536956
Survivin inhibition induces human neural tumor cell death through caspase-independent and -dependent pathways
2001

8/6/19 (Item 19 from file: 5)
0013254012 BIOSIS NO.: 200100425851
Identification of p21 as a target of cycloheximide-mediated facilitation of CD95-mediated apoptosis in human malignant glioma cells
2001

8/6/20 (Item 20 from file: 5)
0013249225 BIOSIS NO.: 200100421064
XIAP : Apoptotic brake and promising therapeutic target
2001

8/6/21 (Item 21 from file: 5)
0013237922 BIOSIS NO.: 200100409761
Gene therapy that inhibits nuclear translocation of nuclear factor kappaB results in tumor necrosis factor alpha-induced apoptosis of human synovial fibroblasts
2000

8/6/22 (Item 22 from file: 5)
0012930183 BIOSIS NO.: 200100102022
Nuclear factor-kappaB-mediated X-linked inhibitor of apoptosis protein expression prevents rat granulosa cells from tumor necrosis factor

alpha-induced apoptosis
2001

8/6/23 (Item 23 from file: 5)
0012919888 BIOSIS NO.: 200100091727
Human ovarian cancer and cisplatin resistance: Possible role of inhibitor
of apoptosis proteins
2001

8/6/24 (Item 24 from file: 5)
0012897193 BIOSIS NO.: 200100069032
Cisplatin (CDDP) sensitizes human osteosarcoma cell to Fas/CD95-mediated
apoptosis by down-regulating FLIP-L expression
2000

8/6/25 (Item 25 from file: 5)
0012810857 BIOSIS NO.: 200000529170
Down-regulation of X-linked inhibitor of apoptosis protein induces
apoptosis in chemoresistant human ovarian cancer cells
2000

8/6/26 (Item 26 from file: 5)
0012731647 BIOSIS NO.: 200000449960
Translational upregulation of X-linked inhibitor of apoptosis (XIAP)
increases resistance to radiation induced cell death
2000

8/6/27 (Item 27 from file: 5)
0012655181 BIOSIS NO.: 200000373494
Metabolic inhibitors sensitize for CD95 (APO-1/Fas)-induced apoptosis by
down-regulating Fas-associated death domain-like interleukin 1-converting
enzyme inhibitory protein expression
2000

8/6/28 (Item 28 from file: 5)
0012599835 BIOSIS NO.: 200000318148
Apoptosis and chemoresistance in human ovarian cancer : Is Xiap a
determinant?
2000

8/6/29 (Item 1 from file: 440)
19534580 Document Delivery Available: 000224692500008 References: 58
TITLE: Loss of XIAP protein expression by RNAi and antisense
approaches sensitizes cancer cells to functionally diverse
chemotherapeutics (ABSTRACT AVAILABLE)
2004
GENUINE ARTICLE#: 865HE

8/6/30 (Item 2 from file: 440)

19532380 Document Delivery Available: 000224732800007 References: 54
**TITLE: Potent antileukemic interactions between flavopiridol and
TRAIL/Apo2L involve flavopiridol-mediated XIAP downregulation (**
ABSTRACT AVAILABLE)
2004
GENUINE ARTICLE#: 865VR

8/6/31 (Item 3 from file: 440)
19532375 Document Delivery Available: 000224732800002 References: 158
**TITLE: Inhibitor of apoptosis proteins: new therapeutic targets in
hematological cancer ? (ABSTRACT AVAILABLE)**
2004
GENUINE ARTICLE#: 865VR

8/6/32 (Item 4 from file: 440)
19524265 Document Delivery Available: 000224756000008 References: 46
**TITLE: IFN-gamma enhances TRAIL-induced apoptosis through IRF-1 (ABSTRACT
AVAILABLE)**
2004
GENUINE ARTICLE#: 866EG

8/6/33 (Item 5 from file: 440)
19519481 Document Delivery Available: 000224701500004 References: 72
**TITLE: The inhibitor of apoptosis protein family and its antagonists in
acute leukemias (ABSTRACT AVAILABLE)**
2004
GENUINE ARTICLE#: 865KL

8/6/34 (Item 6 from file: 440)
19358336 Document Delivery Available: 000224076900013 References: 33
**TITLE: Down-regulation of procaspase-8 expression by focal adhesion kinase
protects HL-60 cells from TRAIL-induced apoptosis (ABSTRACT AVAILABLE)**
2004
GENUINE ARTICLE#: 856WS

8/6/35 (Item 7 from file: 440)
19289502 Document Delivery Available: 000223885100005 References: 28
**TITLE: XIAP and survivin as therapeutic targets for radiation
sensitization in preclinical models of lung cancer (ABSTRACT
AVAILABLE)**
2004
GENUINE ARTICLE#: 854EX

8/6/36 (Item 8 from file: 440)
16717231 Document Delivery Available: 000184578900004 References: 39
**TITLE: Loss of inhibitor of apoptosis proteins as a determinant of
polyamine analog-induced apoptosis in human melanoma cells (ABSTRACT
AVAILABLE)**
2003
GENUINE ARTICLE#: 708QF

8/6/37 (Item 9 from file: 440)
16540070 Document Delivery Available: 000184108700056 References: 40
TITLE: Antisense oligonucleotides targeting XIAP induce apoptosis and enhance chemotherapeutic activity against human lung cancer cells in vitro and in vivo (ABSTRACT AVAILABLE)
2003
GENUINE ARTICLE#: 700JK

8/6/38 (Item 10 from file: 440)
14870510 Document Delivery Available: 000178519100042 References: 44
TITLE: The triterpenoid CDDO induces apoptosis in refractory CLL B cells (ABSTRACT AVAILABLE)
2002
GENUINE ARTICLE#: 602RW

8/6/39 (Item 11 from file: 440)
12401967 References: 47
TITLE: Livin, a novel inhibitor of apoptosis protein family member (ABSTRACT AVAILABLE)
2001
GENUINE ARTICLE#: 398YW

8/6/40 (Item 1 from file: 155)
DIALOG(R)File 155:(c) format only 2004 The Dialog Corp. All rts. reserv.
16255410 PMID: 14990073
[Expression and significance of apoptosis protein inhibitor survivin and XIAP, in patients with myelodysplastic syndromes and in the cell line MUTZ-1]
Jan 2004
Tags: Human; Support, Non-U.S. Gov't
Descriptors: *Apoptosis; *Harringtonines--therapeutic use--TU; *Microtubule-Associated Proteins--genetics--GE; *Myelodysplastic Syndromes--pathology--PA; *Proteins--genetics--GE; Apoptosis--drug effects--DE; Cell Cycle--drug effects--DE; Cell Division--drug effects--DE; Microtubule-Associated Proteins--physiology--PH; Myelodysplastic Syndromes--drug therapy--DT; Oligonucleotides, Antisense--pharmacology--PD; Proteins--physiology--PH; RNA, Messenger--analysis--AN
CAS Registry No.: 0 (Harringtonines); 0 (IAP-like protein, vertebrate); 0 (Microtubule-Associated Proteins); 0 (Oligonucleotides, Antisense); 0 (Proteins); 0 (RNA, Messenger); 0 (survivin); 26833-87-4 (homoharringtonine)

✓ 8/6/41 (Item 2 from file: 155)
DIALOG(R)File 155:(c) format only 2004 The Dialog Corp. All rts. reserv.
14387430 PMID: 10381630
Expression and biological activity of X-linked inhibitor of apoptosis (XIAP) in human malignant glioma.
Apr 1999
Tags: Human; Support, Non-U.S. Gov't
Descriptors: *Apoptosis--genetics--GE; *Brain Neoplasms; *Gene Expression Regulation, Neoplastic; *Glioma; Adenoviridae--genetics--GE; Antigens, CD95--genetics--GE; Antisense Elements (Genetics); Caspases--metabolism--ME;

Gene Transfer Techniques; Neoplasm Proteins--genetics--GE; Proteins
--genetics--GE; Tumor Cells, Cultured--enzymology--EN; Tumor Cells,
Cultured--pathology--PA; Tumor Cells, Cultured--physiology--PH
CAS Registry No.: 0 (Antigens, CD95); 0 (Antisense Elements
(Genetics)); 0 (IAP-like protein, vertebrate); 0 (Neoplasm Proteins); 0
(Proteins)
Enzyme No.: EC 3.4.22.- (Caspases)

8/6/42 (Item 3 from file: 155)
DIALOG(R)File 155:(c) format only 2004 The Dialog Corp. All rts. reserv.

12432968 PMID: 12839953

The histone deacetylase inhibitor MS-275 promotes differentiation or
apoptosis in human leukemia cells through a process regulated by generation
of reactive oxygen species and induction of p21CIP1/WAF1 1.

Jul 1 2003

Tags: Human; Support, Non-U.S. Gov't; Support, U.S. Gov't, P.H.S.
Descriptors: *Apoptosis--drug effects--DE; *Benzamides--pharmacology--PD;
*Cell Differentiation--drug effects--DE; *Cyclins--metabolism--ME; *Histone
Deacetylases--antagonists and inhibitors--AI; *Pyridines--pharmacology--PD;
*Reactive Oxygen Species--metabolism--ME; Cell Cycle--drug effects--DE;
Cell Division--drug effects--DE; HL-60 Cells; Intracellular Membranes--drug
effects--DE; Intracellular Membranes--physiology--PH; K562 Cells; Leukemia
; Membrane Potentials--drug effects--DE; Mitochondria--drug effects--DE;
Mitochondria--physiology--PH; Tumor Cells, Cultured; U937 Cells
CAS Registry No.: 0 (Benzamides); 0 (Cipl protein); 0 (Cyclins); 0
(N-(2-aminophenyl)-4-(N-(pyridin-3-ylmethoxycarbonyl)aminomethyl)benzamide
); 0 (Pyridines); 0 (Reactive Oxygen Species)
Enzyme No.: EC 3.5.1.- (Histone Deacetylases)

8/6/55 (Item 1 from file: 73)
12163838 EMBASE No: 2003265557

The histone deacetylase inhibitor MS-275 promotes differentiation or
apoptosis in human leukemia cells through a process regulated by generation
of reactive oxygen species and induction of p21SUPCIP1/WAF1

01 JUL 2003

8/6/56 (Item 2 from file: 73)
11968278 EMBASE No: 2003076858

Resistance of human ovarian cancer cells to tumor necrosis factor
alpha is a consequence of nuclear factor kappaB-mediated induction of
Fas-associated death domain-like interleukin-1beta-converting enzyme-like
inhibitory protein

01 FEB 2003

8/6/57 (Item 3 from file: 73)
11942842 EMBASE No: 2003053287
Regulation of Apo2L/ tumor necrosis factor-related apoptosis-inducing
ligand-induced apoptosis in thyroid carcinoma cells
2002

8/6/58 (Item 4 from file: 73)
11088683 EMBASE No: 2001106169

Nuclear factor-kappaB-mediated X-linked inhibitor of apoptosis protein expression prevents rat granulosa cells from tumor necrosis factor alpha-induced apoptosis

2001

8/6/59 (Item 1 from file: 144)

DIALOG(R) File 144:(c) 2004 INIST/CNRS. All rts. reserv.

16306294 PASCAL No.: 03-0470411

Antisense oligonucleotides targeting XIAP induce apoptosis and enhance chemotherapeutic activity against human lung cancer cells in vitro and in vivo

2003

English Descriptors: Human; In vitro; Established cell line; Malignant tumor ; Bronchopulmonary; Tumor cell; Biological activity; Animal; Mouse; In vivo; Treatment; Antineoplastic agent; Chemotherapy; Drug combination; Antisense oligonucleotide; Potentiation; Apoptosis; Cell death; Apoptosis inhibitory protein

Broad Descriptors: Rodentia; Mammalia; Vertebrata; Respiratory disease; Lung disease; Bronchus disease; Rodentia; Mammalia; Vertebrata; Appareil respiratoire pathologie; Poumon pathologie; Bronche pathologie; Rodentia; Mammalia; Vertebrata; Aparato respiratorio patologia; Pulmon patologia; Bronquio patologia

French Descriptors: Homme; In vitro; Lignee cellulaire etablie; Tumeur maligne; Bronchopulmonaire; Cellule tumorale; Activite biologique; Animal ; Souris; In vivo; Traitement; Anticancereux; Chimiotherapie; Association medicamenteuse; Oligonucleotide antisens; Potentialisation; Apoptose; Mort cellulaire; Gene XIAP ; Proteine inhibition apoptose

Classification Codes: 002B02R02

Copyright (c) 2003 INIST-CNRS. All rights reserved.

8/6/60 (Item 2 from file: 144)

DIALOG(R) File 144:(c) 2004 INIST/CNRS. All rts. reserv.

15534948 PASCAL No.: 02-0233325

Survivin inhibition induces human neural tumor cell death through caspase-independent and -dependent pathways

2001

English Descriptors: Apoptosis; Tumor cell; Cell death; Cysteine endopeptidases; Caspase; Human

Broad Descriptors: Peptidases; Hydrolases; Enzyme; Peptidases; Hydrolases; Enzyme; Peptidases; Hydrolases; Enzima

French Descriptors: Apoptose; Cellule tumorale; Mort cellulaire; Cysteine endopeptidases; Caspase; Homme; Survivine

Classification Codes: 002B04G01

Copyright (c) 2002 INIST-CNRS. All rights reserved.

8/6/61 (Item 3 from file: 144)

DIALOG(R) File 144: (c) 2004 INIST/CNRS. All rts. reserv.

14624714 PASCAL No.: 00-0295143

Gene therapy that inhibits nuclear translocation of nuclear factor KB results in tumor necrosis factor alpha -induced apoptosis of human synovial fibroblasts
2000

English Descriptors: Rheumatoid arthritis; Treatment; Chemotherapy; Gene therapy; Mechanism of action; Inhibition; Translocation; Apoptosis; Cell death; **Tumor** necrosis factor alpha ; Fibroblast; Synovial membrane; In vitro; Human; Tissue culture; Chronic; Transcription factor NF kappa B
Broad Descriptors: Diseases of the osteoarticular system; Inflammatory joint disease; Immunopathology; Autoimmune disease; Cytokine; Systeme osteoarticulaire pathologie; Rhumatisme inflammatoire; Immunopathologie; Maladie autoimmune; Cytokine; Sistema osteoarticular patologia; Reumatismo inflamatorio; Immunopatologia; Enfermedad autoinmune; Citoquina

French Descriptors: Polyarthrite rhumatoide; Traitement; Chimiotherapie; Therapie genique; Mecanisme action; Inhibition; Translocation; Apoptose; Mort cellulaire; Facteur necrose tumorale alpha ; Fibroblaste; Synoviale ; In vitro; Homme; Culture tissu; Chronique; Facteur transcription NF kappa B

Classification Codes: 002B15D

Copyright (c) 2000 INIST-CNRS. All rights reserved.

?

t sll/medium, k/all

>>>KWIC option is not available in file(s): 399

11/K/1 (Item 1 from file: 5)

DIALOG(R)File 5: Biosis Previews(R)

(c) 2004 BIOSIS. All rts. reserv.

0013914248 BIOSIS NO.: 200200507759

CD40 engagement enhances eosinophil survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

AUTHOR: Bureau Fabrice (Reprint); Seumois Gregory; Jaspar Fabrice; Vanderplasschen Alain; Detry Bruno; Pastoret Paul-Pierre; Louis Renaud; Lekeux Pierre

AUTHOR ADDRESS: Department of Physiology, Faculty of Veterinary Medicine, University of Liege, Sart Tilman, Bat. B42, B-4000, Liege, Belgium** Belgium

JOURNAL: Journal of Allergy and Clinical Immunology 110 (3): p443-449
September, 2002 2002

MEDIUM: print

ISSN: 0091-6749

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

...survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

...ABSTRACT: of the inhibitor of apoptosis protein (IAP) family, namely cellular (c)-IAP1, c-IAP2, and **XIAP**, and 2 antiapoptotic proteins of the Bcl-2 family, namely Bcl-xL and Bfl-1...

...staining with propidium iodide and FITC-conjugated annexin-V. c-IAP2 expression was inhibited with **antisense** oligonucleotides. Results: Freshly isolated eosinophils from healthy and asthmatic patients did not express CD40. Conversely...

...Inhibition of eosinophil apoptosis was accompanied by induction of c-IAP2 but not c-IAP1, **XIAP**, Bcl-xL, or Bfl-1/A1 expression. **Antisense** knockdown of c-iap2 abolished CD40-induced enhancement of eosinophil survival. Sputum cells from asthmatic...

...through induction of c-IAP2 expression and suggest a role for this mechanism in allergic **inflammation**.

DESCRIPTORS:

DISEASES: allergic **inflammation** --

CHEMICALS & BIOCHEMICALS: ...X-inhibitor of apoptosis protein { **XIAP** };

11/K/2 (Item 1 from file: 399)

DIALOG(R)File 399:CA SEARCH(R)

(c) 2004 American Chemical Society. All rts. reserv.

141307497 CA: 141(19)307497m **PATENT**

Use of caspase inhibitors as antiviral agents, and test system for their discovery

INVENTOR(AUTHOR): Ludwig, Stefan; Planz, Oliver; Sedlacek, Hans-Harald; Pleschka, Stephan

LOCATION: Germany,

ASSIGNEE: Medinnova Gesellschaft fur Medizinische Innovationen aus
Akademischer Forschung m.b.H.

PATENT: PCT International ; WO 200485682 A2 DATE: 20041007

APPLICATION: WO 2004DE646 (20040324) *DE 10313636 (20030326)

PAGES: 40 pp. CODEN: PIXXD2 LANGUAGE: German CLASS: C12Q-001/70A

DESIGNATED COUNTRIES: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG; BR; BW; BY;
BZ; CA; CH; CN; CO; CR; CU; CZ; DE; DK; DM; DZ; EC; EE; EG; ES; FI; GB; GD;
GE; GH; GM; HR; HU; ID; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR; LS;
LT; LU; LV; MA; MD; MG; MK; MN; MW; MX; MZ; NA; NI; NO; NZ; OM; PG; PH; PL;
PT; RO; RU; SC; SD; SE; SG; SK; SL; SY; TJ; TM; TN; TR; TT; TZ; UA; UG; US;
UZ; VC; VN; YU; ZA; ZM; ZW DESIGNATED REGIONAL: BW; GH; GM; KE; LS; MW; MZ
; SD; SL; SZ; TZ; UG; ZM; ZW; AM; AZ; BY; KG; KZ; MD; RU; TJ; TM; AT; BE;
BG; CH; CY; CZ; DE; DK; EE; ES; FI; FR; GB; GR; HU; IE; IT; LU; MC; NL; PL;
PT; RO; SE; SI; SK; TR; BF; BJ; CF; CG; CI; CM; GA; GN; GQ; GW; ML; MR; NE;
SN; TD; TG

11/K/3 (Item 1 from file: 144)

DIALOG(R)File 144:Pascal

(c) 2004 INIST/CNRS. All rts. reserv.

14624714 PASCAL No.: 00-0295143

**Gene therapy that inhibits nuclear translocation of nuclear factor KB
results in tumor necrosis factor alpha -induced apoptosis of human
synovial fibroblasts**

ZHANG H G; NING HUANG; DI LIU; BILBAO L; XIAOWU ZHANG; YANG P; TONG ZHOU;
CURIEL D T; MOUNTZ J D

University of Alabama at Birmingham, United States; Gene Therapy Program,
Birmingham, Alabama, United States; University of Alabama at Birmingham,
Birmingham, Alabama, United States; Veterans Administration Medical Center,
Birmingham, Alabama, United States

Journal: Arthritis and rheumatism, 2000, 43 (5) 1094-1105

Language: English

Copyright (c) 2000 INIST-CNRS. All rights reserved.

... factor KB (IKB) dominant-negative adenovirus (AdCMVKB-DN) and an
X-linked inhibitor of apoptosis (**XIAP**) **antisense** adenovirus
(AdCMVXIAP-AS). Primary RA synovial fibroblast (RASf) cell lines were
transfected in vitro, and...

... was no apoptosis after treatment with AdCMV kappa B-DN in the absence
of TNFalpha. **XIAP** is an inhibitor of apoptosis which was up-regulated by
TNFalpha, and this up-regulation...

... greatly enhances apoptosis due to inhibition of an NF-KB-mediated
antiapoptosis signaling pathway, and **XIAP** is a TNFalpha-inducible specific
inhibitor of apoptosis in RA synovial cell lines. This and...

Broad Descriptors: Diseases of the osteoarticular system; **Inflammatory**
joint disease; Immunopathology; Autoimmune disease; Cytokine; Systeme
osteoarticulaire pathologie; Rhumatisme **inflammatoire** ; Immunopathologie
; Maladie autoimmune; Cytokine; Sistema osteoarticular patologia;
Reumatismo inflamatorio; Inmunopatologia; Enfermedad autoinmune;
Citoquina

?

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 15, 2004, 08:23:31 ; Search time 0.001 Seconds
(without alignments)
20.560 Million cell updates/sec

Title: US-10-070-789-8

Sequence: 1 tagactgtccacatttc 20

Scoring table: IDENTITY_NTC
Gapop 10.0 , Gapext 0.5

Searched: 51 segs, 514 residues

Total number of hits satisfying chosen parameters: 102

Minimum DB seq length: 8
Maximum DB seq length: 50

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 53 summaries

Database : fetchg8.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	1	AR103288
2	10	50.0	11	1	AX624749
3	10	50.0	11	1	AX625581
4	10	50.0	11	1	AX632170
5	9.4	47.0	11	1	CO832708
6	8.4	42.0	10	1	AR004345
7	8.4	42.0	10	1	AR004345
8	8.4	42.0	10	1	AR006836
9	8.4	42.0	10	1	AR006836
10	8.4	42.0	10	1	AR227862
11	8.4	42.0	10	1	AR303457
12	8.4	42.0	10	1	AR303598
13	8.4	42.0	10	1	AR303626
14	8.4	42.0	10	1	AX301603
15	8.4	42.0	10	1	BD086244
16	8.4	42.0	10	1	BD166548
17	8.4	42.0	10	1	BD166781
18	8.4	42.0	10	1	AX687084
19	8.4	42.0	10	1	AX687085
20	8.4	42.0	10	1	AX497971
21	8.4	42.0	10	1	AR000247
22	8.4	42.0	10	1	AR150602
23	8.4	42.0	10	1	BD240441
24	8.4	42.0	10	1	BD240561
25	8.4	42.0	10	1	E27852
26	8.4	42.0	10	1	E27855
27	8.4	42.0	10	1	E54835
28	8.4	42.0	10	1	I83580
29	8.4	42.0	10	1	I88948
30	8.4	42.0	10	1	AR200483
31	8.4	42.0	10	1	AR303539
32	8.4	42.0	10	1	AR303605
33	8.4	42.0	10	1	AR371285

C 34	8	40.0	10	1	AR392565	ACCESSION:AR392565
C 35	8	40.0	10	1	AR489506	ACCESSION:AR489506
C 36	8	40.0	10	1	AR491117	ACCESSION:AR491117
C 37	8	40.0	10	1	AX113035	ACCESSION:AX113035
C 38	8	40.0	10	1	AX152807	ACCESSION:AX152807
C 39	8	40.0	10	1	AX152808	ACCESSION:AX152808
C 40	8	40.0	10	1	AX152809	ACCESSION:AX152809
C 41	8	40.0	10	1	AX153055	ACCESSION:AX153055
C 42	8	40.0	10	1	AX153567	ACCESSION:AX153567
C 43	8	40.0	10	1	AX153619	ACCESSION:AX153619
C 44	8	40.0	10	1	AX301314	ACCESSION:AX301314
C 45	8	40.0	10	1	AX301580	ACCESSION:AX301580
C 46	8	40.0	10	1	BD144707	ACCESSION:BD144707
C 47	8	40.0	10	1	BD161205	ACCESSION:BD161205
C 48	8	40.0	10	1	BD161252	ACCESSION:BD161252
C 49	7.4	37.0	9	1	CQ759123	ACCESSION:CQ759123
C 50	7.4	37.0	9	1	AX668629	ACCESSION:AX668629
C 51	7.4	37.0	9	1	AX668630	ACCESSION:AX668630
C 52	7.4	37.0	9	1	AX668813	ACCESSION:AX668813
C 53	7.4	37.0	9	1	AX668814	ACCESSION:AX668814

ALIGNMENTS

RESULT 1	AR103288	Sequence 8 from patent US 6087173.	20 bp	DNA	linear	PAT 14-FEB-2001
LOCUS	AR103288	Sequence 8 from patent US 6087173.	20 bp	DNA	linear	PAT 14-FEB-2001
DEFINITION	AR103288	Sequence 8 from patent US 6087173.	20 bp	DNA	linear	PAT 14-FEB-2001
ACCESSION	AR103288	Sequence 8 from patent US 6087173.	20 bp	DNA	linear	PAT 14-FEB-2001
VERSION	AR103288.1	GI:12814876	20 bp	DNA	linear	PAT 14-FEB-2001
KEYWORDS	Unknown.					
SOURCE	Unknown.					
ORGANISM	Unknown.					
REFERENCE	1 (bases 1 to 20)					
AUTHORS	Bennett,C.Frank., Ackermann,E.J. and Cowsett,L.M.					
TITLE	Antisense modulation of X-linked inhibitor of apoptosis expression					
JOURNAL	Patent: US 6087173-A 8 11-JUL-2000;					
FEATURES	location/Qualifiers					
source	1..20					
	/organism="unknown"					
	/mol_type="unassigned DNA"					
Query Match	100.0%; Score 20; DB 1; Length 20;					
Best Local Similarity	100.0%; Pred. No. 0.049;					
Matches	20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;					
QY	1 TAGGACTGTGCACCTTTTC 20					
DB	1 TAGGACTGTGCACCTTTTC 20					
RESULT 2	AX624749/c					
LOCUS	AX624749	Sequence 11 bp	DNA	linear		PAT 21-FEB-2003
DEFINITION	AX624749	Sequence 11 bp	DNA	linear		PAT 21-FEB-2003
ACCESSION	AX624749	Sequence 11 bp	DNA	linear		PAT 21-FEB-2003
VERSION	AX624749.1	GI:28452690				
KEYWORDS						
SOURCE	Homo sapiens (human)					
ORGANISM	Homo sapiens					
REFERENCE	Eukariota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;					
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.					
TITLE	Peterohn,D., Conradt,M. and Hofmann,K.					
JOURNAL	Method for determining homeostasis of the skin					
FEATURES	Patent: WO 02053774-A 1790 11-JUL-2002;					
source	Henkel Kommandgesellschaft auf Aktien (DE)					
	Location/Qualifiers					
	1..11					
	/organism="Homo sapiens"					
	/mol_type="unassigned DNA"					

OTHER INFORMATION: Description of Artificial Sequence: Zfp 1 target sequence

Query Match 32.0%; Score 6.4; DB 1; Length 8;
 Best Local Similarity 87.5%; Pred. No. 0;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 11 CCACCTT 18
 |||||
 8 CCACCTT 1

DB

RESULT 3
 CF339699/c 8 bp mRNA linear EST 18-AUG-2003
 LOCUS RCL1--05-K22.g1 Regenerated callus lambda phage cDNA library (RCL1)
 DEFINITION Oryza sativa (japonica cultivar-group) cDNA clone RCL1--05-K22,
 mRNA sequence.

ACCESSION CF339699.1 GI:33827769
 VERSION EST.
 KEYWORDS Oryza sativa (japonica cultivar-group)
 SOURCE Oryza sativa (japonica cultivar-group)
 ORGANISM Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoideae; Oryzaceae; Oryza.

REFERENCE 1 (bases 1 to 8)
 Kim,J.S., Jun,K.M., Cheong,P.J., Kim,M.J., Lee,T.H., Shin,Y.C.,
 Song,S.I., Kim,J.K., Kim,Y.-K. and Nahm,B.H.
 Large-scale Sequencing Analysis of Rice ESTs
 Unpublished (2003)
 CONTACT: Nahm B.H.
 GENOMICS AND GENETICS INSTITUTE, GREENGENE BIOTECH INC., DIVISION
 OF BIOLOGICAL AND BIOINFORMATICS, MYONGJI UNIVERSITY
 YONGIN, KYEONGGI, KOREA
 TEL: 82 31 330 6193
 FAX: 82 31 321 6355
 EMAIL: bhnahm@gbio.com, bhnahm@bio.myongji.ac.kr.

FEATURES
 source location/Qualifiers

1..8
 /organism="Oryza sativa (japonica cultivar-group)"
 /mol_type="mRNA"
 /cultivar="Nackdong"
 /db_xref="taxon:39947"
 /clone="RCL1--05-K22"
 /tissue_type="callus"
 /dev_stage="proliferated callus on 2N6 media for 30 days"
 /lab_host="E.coli SOLR"
 /clone_lib="Regenerated callus lambda phage cDNA library (RCL1)"
 /note="Vector: pBluescript SK(+); Site_1: SctI; Site_2: XhoI; cDNA was inserted into lambda Uni-ZAP XR vector at 5' end with SctI and 3' end with XhoI site. Callus was induced on 2N6 media for 30 days and cultured for 36hrs on regenerated media"

Query Match 32.0%; Score 6.4; DB 1; Length 8;
 Best Local Similarity 87.5%; Pred. No. 0;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 11 CCACCTT 18
 |||||
 8 CCACCTT 1

DB

RESULT 4
 CF340204/c 8 bp mRNA linear EST 18-AUG-2003
 LOCUS RCL1--07-E15.g1 Regenerated callus lambda phage cDNA library (RCL1)
 DEFINITION Oryza sativa (japonica cultivar-group) cDNA clone RCL1--07-E15,
 mRNA sequence.

ACCESSION CF340204.1 GI:33828768
 VERSION EST.
 KEYWORDS Oryza sativa (japonica cultivar-group)
 SOURCE Oryza sativa (japonica cultivar-group)
 ORGANISM

REFERENCE 1 (bases 1 to 8)
 Kim,J.S., Jun,K.M., Cheong,P.J., Kim,M.J., Lee,T.H., Shin,Y.C.,
 Song,S.I., Kim,J.K., Kim,Y.-K. and Nahm,B.H.
 Large-scale Sequencing Analysis of Rice ESTs
 Unpublished (2003)
 CONTACT: Nahm B.H.
 GENOMICS AND GENETICS INSTITUTE, GREENGENE BIOTECH INC., DIVISION
 OF BIOLOGICAL AND BIOINFORMATICS, MYONGJI UNIVERSITY
 YONGIN, KYEONGGI, KOREA
 TEL: 82 31 330 6193
 FAX: 82 31 321 6355
 EMAIL: bhnahm@gbio.com, bhnahm@bio.myongji.ac.kr.

FEATURES
 source location/Qualifiers

1..8
 /organism="Oryza sativa (japonica cultivar-group)"
 /mol_type="mRNA"
 /cultivar="Nackdong"
 /db_xref="taxon:39947"
 /clone="RCL1--07-E15"
 /tissue_type="callus"
 /dev_stage="proliferated callus on 2N6 media for 30 days"
 /lab_host="E.coli SOLR"
 /clone_lib="Regenerated callus lambda phage cDNA library (RCL1)"
 /note="Vector: pBluescript SK(+); Site_1: SctI; Site_2: XhoI; cDNA was inserted into lambda Uni-ZAP XR vector at 5' end with SctI and 3' end with XhoI site. Callus was induced on 2N6 media for 30 days and cultured for 36hrs on regenerated media"

Query Match 32.0%; Score 6.4; DB 1; Length 8;
 Best Local Similarity 87.5%; Pred. No. 0;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 11 CCACCTT 18
 |||||
 8 CCACCTT 1

DB

Search completed: November 15, 2004, 08:31:01
 Job time : 0.001 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OW nucleic - nucleic search, using sw model

Run on: November 15, 2004, 08:27:34 ; Search time 0.001 Seconds
(without alignments)
14.400 Million cell updates/sec

Title: US-10-070-789-8
Percent score: 20
Sequence: 1 taggactgtccacatttc 20

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 0.5

Searched: 37 seqs, 360 residues

Total number of hits satisfying chosen parameters: 74

Minimum DB seq length: 8
Maximum DB seq length: 50

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 39 summaries

Database : fetchrn18.seq.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	1	US-09-392-580-8
2	8.4	42.0	10	1	US-08-460-806-27
3	8.4	42.0	10	1	US-08-460-806-27
4	8.4	42.0	10	1	US-08-325-630-27
5	8.4	42.0	10	1	US-08-325-630-27
6	8.4	42.0	10	1	US-08-388-353-191
7	8.4	42.0	10	1	US-08-388-353-191
8	8.4	42.0	10	1	US-08-488-551B-191
9	8.4	42.0	10	1	US-08-488-551B-191
10	8.4	42.0	10	1	US-08-488-551B-191
11	8.4	42.0	10	1	US-08-994-035C-1
12	8.4	42.0	10	1	US-09-395-861-1
13	8.4	42.0	10	1	US-09-508-753B-182
14	8.4	42.0	10	1	US-09-508-753B-182
15	8.4	42.0	10	1	US-09-508-753B-351
16	8.4	42.0	8	1	US-08-859-954-208
17	8.4	42.0	8	1	US-08-859-954-230
18	8.4	42.0	8	1	US-08-859-954-540
19	8.4	42.0	10	1	US-08-866-116A-45
20	8.4	42.0	10	1	US-08-866-116A-45
21	8.4	42.0	10	1	US-08-847-108-45
22	8.4	42.0	10	1	US-08-866-113A-52
23	8.4	42.0	10	1	US-08-847-095A-45
24	8.4	42.0	10	1	US-08-866-113A-52
25	8.4	42.0	10	1	US-08-150-156A-26
26	8.4	42.0	10	1	US-08-150-156A-26
27	8.4	42.0	10	1	US-08-686-114B-52
28	8.4	42.0	10	1	US-08-686-114B-52
29	8.4	42.0	10	1	US-08-508-753B-264
30	8.4	42.0	10	1	US-08-508-753B-330
31	8.4	42.0	10	1	US-08-337-304-52
32	8.4	42.0	10	1	US-08-462-977B-22
33	7	35.0	8	1	US-08-662-963-3

34	7	35.0	8	1	US-08-662-963-10	Sequence 10, Appl
35	7	35.0	8	1	US-08-662-963-16	Sequence 16, Appl
36	7	35.0	8	1	US-08-662-963-19	Sequence 19, Appl
37	7	35.0	8	1	US-08-859-954-125	Sequence 125, Appl
38	7	35.0	8	1	US-08-859-954-245	Sequence 245, Appl
39	7	35.0	8	1	US-08-859-954-384	Sequence 384, Appl

ALIGNMENTS

```
RESULT 1
US-09-392-580-8
Sequence 8, Application US/09392580
Patent No. 6087173
GENERAL INFORMATION:
APPLICANT: C. Frank Bennett
APPLICANT: Elizabeth J. Ackermann
APPLICANT: Lex M. Cosset
TITLE OF INVENTION: ANTISENSE MODULATION OF X-LINKED INHIBITOR OF APOPTOSIS EXPRESSIC
FILE REFERENCE: RFS-0072
CURRENT APPLICATION NUMBER: US/09/392,580
CURRENT FILING DATE: 1999-09-05
NUMBER OF SEQ ID NOS: 47
SEQ ID NO 8
LENGTH: 20
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Antisense Oligonucleotide
US-09-392-580-8

Query Match      100.0%: Score 20; DB 1; Length 20;
Best local Similarity 100.0%: Pred. No. 0.044;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TAGGACTGTCCACCTTTTC 20
Db 1 TAGGACTGTCCACCTTTTC 20

RESULT 2
US-08-460-806-27
Sequence 27, Application US/08460806
Patent No. 5747241
GENERAL INFORMATION:
APPLICANT: MIYAMURA, TATSUO
APPLICANT: SAITO, IZUMU
APPLICANT: HARADA, SHIZUKO
APPLICANT: HONDA, YOSHIKAZU
TITLE OF INVENTION: DIAGNOSTIC REAGENT FOR HEPATITIS C
NUMBER OF SEQUENCES: 28
CORRESPONDENCE ADDRESS:
ADDRESSEE: OHLON, SPIVAK, MCLELLAND, MAIER & NEUSTADT,
STREET: 1755 S. Jefferson Davis Highway, Suite 400
CITY: Arlington
STATE: Virginia
COUNTRY: U.S.A.
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/460,806
FILING DATE: 02-JUN-1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/325,630
FILING DATE: 19-OCT-1994
APPLICATION NUMBER: US 07/956,993
```

Set	Items	Description
S1	6121	(X-LINKED INHIBITOR OF APOPTOSIS) OR XIAP OR HIAP-1 OR HILP OR (X-LINKED IAP) OR MIHA
S2	257315	ANTISENSE OR RIBOZYME OR TRIPLEX OR SIRNA OR RNAI OR (SHORT INTERFER? RNA) OR (RNA INTERFER?)
S3	10136532	CANCER? OR TUMOR? OR TUMOUR? OR CARCINOMA?
S4	1886463	INFLAMMATORY OR INFLAMMATION
S5	1147	(FOLLICULAR ATRESIA?) OR (ATRESIA? FOLLICULAR)
S6	515	S1 AND S2
S7	436	S6 AND S3
S8	3487542	6 AND S3 OR S4 OR S5
S9	1888026	S6 AND S3 OR S4 OR S5
S10	26	S6 AND S4
S11	2	S6 AND S5
S12	0	9 RD
S13	3487542	6 AND S3 OR S4 OR S5
S14	1888026	S6 AND S3 OR S4 OR S5
S15	436	S6 AND S3
S16	165	RD (unique items)
S17	10	S16 NOT PY>2000

JOURNAL REGION: USA
JOURNAL SUBJECT: Business; General News
? S S16 NOT PY>2000
Processing
Processing
Processed 10 of 18 files ...
Processing
>>>One or more prefixes are unsupported
>>> or undefined in one or more files.
Processing
Processing
Completed processing all files
165 S16
94675125 PY>2000
S17 10 S16 NOT PY>2000
? T S17/MEDIUM,K/ALL
>>>KWIC option is not available in file(s): 398, 399

17/K/1 (Item 1 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.

0013237922 BIOSIS NO.: 200100409761

Gene therapy that inhibits nuclear translocation of nuclear factor kappaB results in tumor necrosis factor alpha-induced apoptosis of human synovial fibroblasts

AUTHOR: Zhang Huang-Ge; Huang Ning; Liu Di; Bilbao Lupita; Zhang Xiaowu; Yang Pingar; Zhou Tong; Curiel David T; Mountz John D (Reprint)
AUTHOR ADDRESS: Department of Medicine, Division of Clinical Immunology and Rheumatology, The University of Alabama at Birmingham, 701 South 19th Street, LHRB 473, Birmingham, AL, 35294, USA**USA
JOURNAL: Arthritis and Rheumatism 43 (5): p1094-1105 May, 2000 2000
MEDIUM: print
ISSN: 0004-3591
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English

Gene therapy that inhibits nuclear translocation of nuclear factor kappaB results in tumor necrosis factor alpha-induced apoptosis of human synovial fibroblasts

ABSTRACT: Objective. **Tumor** necrosis factor alpha (TNFalpha) increases the survival and proliferation of human rheumatoid arthritis (RA) cell...

...factor kappaB (IkappaB) dominant-negative adenovirus (AdCMVikappaB-DN) and an X-linked inhibitor of apoptosis (**XIAP**) **antisense** adenovirus (AdCMVXIAP-AS). Primary RA synovial fibroblast (RASf) cell lines were transfected in vitro, and...

...TNFalpha. There was no apoptosis after treatment with AdCMVikappaB-DN in the absence of TNFalpha. **XIAP** is an inhibitor of apoptosis which was up-regulated by TNFalpha, and this up-regulation...

...greatly enhances apoptosis due to inhibition of an NF-kappaB-mediated antiapoptosis signaling pathway, and **XIAP** is a TNFalpha-inducible specific inhibitor of apoptosis in RA synovial cell lines. This and...

DESCRIPTORS:

CHEMICALS & BIOCHEMICALS: ... tumor necrosis factor-alpha

17/K/2 (Item 2 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.

0012897193 BIOSIS NO.: 200100069032
Cisplatin (CDDP) sensitizes human osteosarcoma cell to Fas/CD95-mediated apoptosis by down-regulating FLIP-L expression
AUTHOR: Kinoshita Hirokazu; Yoshikawa Hideshi; Shiki Kazuhiko; Hamada Yoshiki; Nakajima Yasuo; Tasaka Kachio (Reprint)
AUTHOR ADDRESS: Department of Parasitology and Immunology, Yamanashi Medical University, 1110 Shimokato, Tamaho-cho, Yamanashi, 409-3898, Japan**Japan
JOURNAL: International Journal of Cancer 88 (6): p986-991 15 December, 2000 2000
MEDIUM: print
ISSN: 0020-7136
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English

...ABSTRACT: escape from Fas/CD95-mediated apoptosis induced by immunosurveillance(NK cells and T cells) in tumor cells are correlated to **tumorigenicity**. Human osteosarcoma cell MG-63 constitutively expressed cell surface Fas antigen but was resistant to...
...form(FLIP-L), which was a novel anti-apoptotic protein and had a potency of **tumorigenicity**. CDDP down-regulated FLIP-L in a time-dependent manner in MG-63 cells but did not influence expression of other anti-apoptotic molecules such as **XIAP**, c-IAP-1, c-IAP-2, FADD or pro-caspase-8. Moreover, **antisense** oligonucleotide to FLIP-L confirmed that down-regulation of FLIP-L induced sensitization to Fas...

DESCRIPTORS:

...MAJOR CONCEPTS: Tumor Biology
CHEMICALS & BIOCHEMICALS: ... **XIAP** ;

17/K/3 (Item 3 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.

0012810857 BIOSIS NO.: 200000529170
Down-regulation of X-linked inhibitor of apoptosis protein induces apoptosis in chemoresistant human ovarian cancer cells
AUTHOR: Sasaki Hiromasa; Sheng YingLun; Kotsuji Fumikazu; Tsang Benjamin K (Reprint)
AUTHOR ADDRESS: Loeb Health Research Institute, 725 Parkdale Avenue, Ottawa, Ontario, K1Y 4E9: btsang@lri.ca, Canada**Canada
JOURNAL: Cancer Research 60 (20): p5659-5666 October 15, 2000 2000
MEDIUM: print
ISSN: 0008-5472
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English

Down-regulation of X-linked inhibitor of apoptosis protein induces apoptosis in chemoresistant human ovarian cancer cells

ABSTRACT: Cisplatin-centered chemotherapy is a key treatment for ovarian **cancer**, but resistance to chemotherapeutic agents remains a major cause of treatment failure. Multiple factors are...
 ...of this chemoresistance. Although it has been demonstrated that X-linked inhibitor of apoptosis protein (**Xiap**) prevents apoptosis by inhibiting effector caspases, if and how it is important in chemoresistance in ovarian **cancer** has not been studied. The effects of **Xiap** down-regulation and/or restoration of wild type p53 by recombinant adenovirus infection were examined on four ovarian epithelial **cancer** cell lines (C13*, A2780-s (wild type p53), A2780-cp (mutant p53), and SKOV3 (null p53)). Apoptosis and protein expression (e.g., **Xiap**, caspase-3, p53, MDM2, and p21waf1) were assessed by Hoechst 33258 stain and Western blot, respectively. We demonstrated that **Xiap** down-regulation following adenoviral **antisense** expression induces apoptosis in the wild-type p53 cells, but not in the mutated or null cells. **Xiap** down-regulation resulted in caspase-3 activation, caspase-mediated MDM2 processing, and p53 accumulation. Restoration...
 ...type p53 in the p53-mutated or -null cells significantly enhanced the proapoptotic effect of **Xiap antisense** expression. Down-regulation of **Xiap** induced apoptosis in chemoresistant ovarian **cancer** cells, a process dependent on p53 status.

DESCRIPTORS:

...MAJOR CONCEPTS: **Tumor** Biology
 ...ORGANISMS: chemoresistance, human ovarian **cancer** cell line, in-vitro model system, mutant p53 gene expression...
 ...chemoresistance, human ovarian **cancer** cell line, in-vitro model system, wild type p53 gene expression...
 ...chemoresistance, human ovarian **cancer** cell line, in-vitro model system, wild type p53 gene expression...
 ...chemoresistance, human ovarian **cancer** cell line, in-vitro model system, null p53 gene expression
 CHEMICALS & BIOCHEMICALS: ...X-linked inhibitor of apoptosis downregulation role, caspase-mediated processing, **tumor** cell expression...
 ... **antisense** transfer-mediated downregulation, chemoresistant **tumor** cell apoptosis induction, recombinant adenovirus-mediated **antisense** transfer, **tumor** cell expression...
 ...X-linked inhibitor of apoptosis protein-induced **tumor** cell expression
 ...
 ...antineoplastic-drug, **tumor** cell resistance...
 ...X-linked inhibitor of apoptosis downregulation role, **tumor** cell accumulation

17/K/4 (Item 4 from file: 5)

DIALOG(R) File 5: Biosis Previews(R)

(c) 2004 BIOSIS. All rts. reserv.

0012731647 BIOSIS NO.: 200000449960

**Translational upregulation of X-linked inhibitor of apoptosis (XIAP)
increases resistance to radiation induced cell death**

AUTHOR: Holcik Martin; Yeh Chiaoli; Korneluk Robert G (Reprint); Chow Terry
AUTHOR ADDRESS: Molecular Genetics, Research Institute, Children's Hospital
of Eastern Ontario, 401 Smyth Road, Ottawa, Ontario, K1H 8L1, Canada**
Canada

JOURNAL: Oncogene 19 (36): p4174-4177 24 August, 2000 2000

MEDIUM: print

ISSN: 0950-9232

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

**Translational upregulation of X-linked inhibitor of apoptosis (XIAP)
increases resistance to radiation induced cell death**

ABSTRACT: Inhibitory regulators of apoptosis play a critical role in the responsiveness of **tumour** cells to cytotoxic agents. The X-linked inhibitor of apoptosis protein (**XIAP**) is a member of a novel family of Inhibitor of Apoptosis (IAP) proteins. Here we show that acute low dose ionizing irradiation results in the translational upregulation of **XIAP** that correlates with an increased resistance to radiation in non-small cell lung **carcinoma** . This upregulation is mediated by an internal ribosome binding mechanism via an IRES element located within a **XIAP** 5' UTR. Transient overexpression of **XIAP** rendered human **carcinoma** cells resistant to low dose gamma-irradiation. By contrast, the **antisense** targeting of **XIAP** resulted in increased cell death following irradiation advocating a distinct role for **XIAP** in radiation resistant phenotype of human **cancers** .

DESCRIPTORS:

...MAJOR CONCEPTS: **Tumor** Biology

DISEASES: **cancer** --...

...non-small cell lung **carcinoma** --

...MESH TERMS: **Carcinoma** , Non-Small-Cell Lung (MeSH

CHEMICALS & BIOCHEMICALS: ... **X-linked inhibitor of apoptosis** ...

... **antisense** targeting, translation upregulation...

... **tumor** cell responsiveness

17/K/5 (Item 5 from file: 5)

DIALOG(R)File 5: Biosis Previews(R)

(c) 2004 BIOSIS. All rts. reserv.

0012655181 BIOSIS NO.: 200000373494

**Metabolic inhibitors sensitize for CD95 (APO-1/Fas)-induced apoptosis by
down-regulating Fas-associated death domain-like interleukin 1-converting
enzyme inhibitory protein expression**

AUTHOR: Fulda Simone; Meyer Eric; Debatin Klaus-Michael (Reprint)

AUTHOR ADDRESS: University Children's Hospital, Prittwitzstrasse 43,
D-89075, Ulm, Germany**Germany

JOURNAL: Cancer Research 60 (14): p3947-3956 July 15, 2000 2000

MEDIUM: print

ISSN: 0008-5472

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

...ABSTRACT: both molecules have a high turnover rate. Selective down-regulation of FLIP expression by FLIP **antisense** oligonucleotides sensitized for CD95-induced apoptosis. Reduction of FLIP levels resulted in undetectable amounts of...

...sensitized for subsequent CD95 stimulation compared with cells without pretreatment. CHX or ActD also reduced **XIAP** expression and similarly sensitized for **tumor** necrosis factor-related apoptosis-inducing ligand- or **tumor** necrosis factor-alpha-induced apoptosis. Because blockade of death receptor triggering by FLIP overexpression has recently been implicated in **tumorigenesis** and treatment resistance in vivo, strategies to inhibit FLIP expression, e.g., by metabolic inhibitors, may prove to be a useful complementary tool for the treatment of **cancer**.

DESCRIPTORS:

...MAJOR CONCEPTS: **Tumor** Biology

CHEMICALS & BIOCHEMICALS: ...metabolic inhibitor drug sensitization, **tumor** cell apoptosis induction...

...metabolic inhibitor drug-induced downregulation, **tumor** cell, expression...

17/K/6 (Item 6 from file: 5)

DIALOG(R)File 5:Biosis-Previews(R)

(c) 2004 BIOSIS. All rts. reserv.

0012599835 BIOSIS NO.: 200000318148

Apoptosis and chemoresistance in human ovarian cancer : Is Xiap a determinant?

AUTHOR: Li Julang; Sasaki Hiromasa; Sheng Ying Lun; Schneiderman Danielle; Xiao Chao Wu; Kotsuji Fumikazu; Tsang Benjamin K (Reprint)

AUTHOR ADDRESS: Department of Obstetrics and Gynaecology, Ottawa Hospital, 1053 Carling Avenue, Civic Campus, Ottawa, ON, K1Y 4E9, Canada**Canada

JOURNAL: Biological Signals and Receptors 9 (2): p122-130 March-April, 2000 2000

MEDIUM: print

ISSN: 1422-4933

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

Apoptosis and chemoresistance in human ovarian cancer : Is Xiap a determinant?

ABSTRACT: Cisplatin-induced apoptosis in epithelial ovarian **cancer** cells is in part a consequence of suppressed **Xiap** expression and upregulation of the Fas/FasL system. Changes in the expression of these 'cell...

...genes lead to activation of caspase-3, and cleavage of MDM2 and FAK.

Failure of **cancer** cells to maintain a balance in the expression of these genes in favor of apoptotic cell death may be an important factor of chemoresistance. **Xiap** may be a novel target for gene therapy of human ovarian epithelial **cancer** and, dependent on P53 status, expression of **Xiap antisense** alone or in combination with wild-type P53 sense may offer a new approach for the treatment of the chemoresistant **cancer**.

DESCRIPTORS:

DISEASES: ovarian cancer --

17/K/7 (Item 1 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

14387430 PMID: 10381630

Expression and biological activity of X-linked inhibitor of apoptosis (XIAP) in human malignant glioma.

Wagenknecht B; Glaser T; Naumann U; Kugler S; Isenmann S; Bahr M; Korneluk R; Liston P; Weller M

Laboratory of Molecular Neuro-Oncology, Hoppe-Seyler-Strasse 3, Germany.

Cell death and differentiation (ENGLAND) Apr 1999, 6 (4) p370-6,

ISSN 1350-9047 Journal Code: 9437445

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Expression and biological activity of X-linked inhibitor of apoptosis (XIAP) in human malignant glioma.

... via direct inhibition of caspases. Here, we report that human malignant glioma cell lines express **XIAP**, HIAP-1 and HIAP-2 mRNA and proteins. NAIP was not expressed. IAP proteins were...

... exposure to CD95L and a protein synthesis inhibitor, CHX, to acquire sensitivity to apoptosis. Adenoviral **XIAP** gene transfer blocked caspase 8 and 3 processing in both cell lines in the absence of CHX. Apoptosis was blocked in the absence and in the presence of CHX. However, **XIAP** failed to block caspase 8 processing in LN-229 cells in the presence of CHX. There was considerable overlap of the effects of **XIAP** on caspase processing with those of BCL-2 and the viral caspase inhibitor crm-A...

... of these cells to apoptotic stimuli that directly target caspases, including radiochemotherapy and immune-mediated **tumor** cell lysis.

; Adenoviridae--genetics--GE; Antigens, CD95--genetics--GE; **Antisense** Elements (Genetics); Caspases--metabolism--ME; Gene Transfer Techniques; Neoplasm Proteins--genetics--GE; Proteins--genetics--GE; **Tumor** Cells, Cultured--enzymology--EN; **Tumor** Cells, Cultured--pathology--PA; **Tumor** Cells, Cultured--physiology--PH

Chemical Name: Antigens, CD95; **Antisense** Elements (Genetics); IAP-like protein, vertebrate; Neoplasm Proteins; Proteins; Caspases

17/K/8 (Item 1 from file: 399)

DIALOG(R)File 399:CA SEARCH(R)

(c) 2004 American Chemical Society..All rts. reserv.

130137344 CA: 130(11)137344n PATENT

XAF proteins interacting with IAPs (inhibitors of apoptosis proteins) and the genes encoding them and their use in the therapeutic regulation of apoptosis

INVENTOR(AUTHOR): Korneluk, Robert; Tamai, Katsuyuki; Liston, Peter; Mackenzie, Alexander E.; Baird, Stephen

LOCATION: Can.,

ASSIGNEE: University of Ottawa

PATENT: European Pat. Appl. ; EP 892048 A2 DATE: 19990120
APPLICATION: EP 98113003 (19980713) *US 52402 (19970714) *US 54491
(19970801) *US 56338 (19970818)
PAGES: 101 pp. CODEN: EPXXDW LANGUAGE: English CLASS: C12N-015/12A;
C12N-015/11B; C07K-014/47B; C12N-015/85B; C12N-005/10B; A01K-067/027B;
G01N-033/50B; A61K-038/17B; A61K-048/00B; C12Q-001/68B; C12P-021/08B
DESIGNATED COUNTRIES: AT; BE; CH; DE; DK; ES; FR; GB; GR; IT; LI; LU; NL;
SE; MC; PT; IE; SI; LT; LV; FI; RO

17/K/9 (Item 2 from file: 399)
DIALOG(R) File 399:CA SEARCH(R)
(c) 2004 American Chemical Society. All rts. reserv.

129197985 CA: 129(16)197985m PATENT
Detection and modulation of protein inhibitors of apoptosis (IAPs) and neuronal apoptosis-inhibiting protein (NAIP) for the diagnosis and treatment of proliferative disease
INVENTOR(AUTHOR): Korneluk, Robert; MacKenzie, Alexander E.; Liston, Peter; Baird, Stephen; Tsang, Benjamin; Pratt, Christine
LOCATION: Can.,
ASSIGNEE: University of Ottawa
PATENT: PCT International ; WO 9835693 A2 DATE: 19980820
APPLICATION: WO 98IB781 (19980213) *US 800929 (19970213)
PAGES: 148 pp. CODEN: PIXXD2 LANGUAGE: English CLASS: A61K-038/17A;
A61K-031/70B; A61K-039/395B; C12N-015/11B; C12Q-001/68B; G01N-033/50B;
G01N-033/574B; A01K-067/027B; C12N-015/00B DESIGNATED COUNTRIES: AL; AM;
AT; AU; AZ; BA; BB; BG; BR; BY; CA; CH; CN; CU; CZ; DE; DK; EE; ES; FI; GB;
GE; GH; GM; GW; HU; ID; IL; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR; LS; LT;
LU; LV; MD; MG; MK; MN; MW; MX; NO; NZ; PL; PT; RO; RU; SD; SE; SG; SI; SK;
SL; TJ; TM; TR; TT; UA; UG; US; UZ; VN; YU; ZW; AM; AZ; BY; KG; KZ; MD; RU;
TJ; TM DESIGNATED REGIONAL: GH; GM; KE; LS; MW; SD; SZ; UG; ZW; AT; BE; CH;
; DE; DK; ES; FI; FR; GB; GR; IE; IT; LU; MC; NL; PT; SE; BF; BJ; CF; CG;
CI; CM; GA; GN; ML; MR; NE; SN; TD; TG

17/K/10 (Item 1 from file: 144)
DIALOG(R) File 144:Pascal
(c) 2004 INIST/CNRS. All rts. reserv.

14624714 PASCAL No.: 00-0295143
Gene therapy that inhibits nuclear translocation of nuclear factor KB results in tumor necrosis factor alpha -induced apoptosis of human synovial fibroblasts
ZHANG H G; NING HUANG; DI LIU; BILBAO L; XIAOWU ZHANG; YANG P; TONG ZHOU;
CURIEL D T; MOUNTZ J D
University of Alabama at Birmingham, United States; Gene Therapy Program,
Birmingham, Alabama, United States; University of Alabama at Birmingham,
Birmingham, Alabama, United States; Veterans Administration Medical Center,
Birmingham, Alabama, United States
Journal: Arthritis and rheumatism, 2000, 43 (5) 1094-1105
Language: English

Copyright (c) 2000 INIST-CNRS. All rights reserved.

Gene therapy that inhibits nuclear translocation of nuclear factor KB results in tumor necrosis factor alpha -induced apoptosis of human synovial fibroblasts

Objective. **Tumor** necrosis factor α (TNF α) increases the survival and proliferation of human rheumatoid arthritis (RA) cell...
... factor KB (IKB) dominant-negative adenovirus (AdCMVIKB-DN) and an X-linked inhibitor of apoptosis (**XIAP**) **antisense** adenovirus (AdCMVXIAP-AS). Primary RA synovial fibroblast (RASf) cell lines were transfected in vitro, and...

... was no apoptosis after treatment with AdCMVI kappa B-DN in the absence of TNF α . **XIAP** is an inhibitor of apoptosis which was up-regulated by TNF α , and this up-regulation...

... greatly enhances apoptosis due to inhibition of an NF-KB-mediated antiapoptosis signaling pathway, and **XIAP** is a TNF α -inducible specific inhibitor of apoptosis in RA synovial cell lines. This and...

English Descriptors: Rheumatoid arthritis; Treatment; Chemotherapy; Gene therapy; Mechanism of action; Inhibition; Translocation; Apoptosis; Cell death; **Tumor** necrosis factor α ; Fibroblast; Synovial membrane; In vitro; Human; Tissue culture; Chronic; Transcription factor NF...

French Descriptors: Polyarthrite rhumatoïde; Traitement; Chimiothérapie; Thérapie génique; Mécanisme d'action; Inhibition; Translocation; Apoptose; Mort cellulaire; Facteur de nécrose **tumorale** α ; Fibroblaste; Synoviale; In vitro; Homme; Culture tissulaire; Chronique; Facteur de transcription NF kappa B

Spanish Descriptors: Poliartritis reumatoidea; Tratamiento; Quimioterapia; Terapia génica; Mecanismo de acción; Inhibición; Translocación; Apoptosis; Muerte celular; Factor de necrosis **tumoral** α ; Fibroblasto; Sinovial; In vitro; Hombre; Cultivo tejido; Crónico

?

18	6121	(X-LINKED INHIBITOR OF APOPTOSIS) OR XIAP OR HIAP-1 OR HILP OR (X-LINKED IAP) OR MIHA
S19	257315	ANTISENSE OR RIBOZYME OR TRIPLEX OR SIRNA OR RNAI OR (SHORT INTERFER? RNA) OR (RNA INTERFER?)
S20	10136532	CANCER? OR TUMOR? OR TUMOUR? OR CARCINOMA?
S21	1886463	INFLAMMATORY OR INFLAMMATION
S22	1147	(FOLLICULAR ATRESIA?) OR (ATRESIA? FOLLICULAR)
S23	515	S18 AND S19
S24	436	S23 AND S20
S25	3487542	6 AND S20 OR S21 OR S22
S26	1888026	S23 AND S20 OR S21 OR S22
S27	26	S23 AND S21
S28	2	S23 AND S22
S29	0	9 RD
S30	3487542	6 AND S20 OR S21 OR S22
S31	1888026	S23 AND S20 OR S21 OR S22
S32	436	S23 AND S20
S33	165	RD (unique items)
S34	10	S33 NOT PY>2000

Set	Items	Description
S1	6121	(X-LINKED INHIBITOR OF APOPTOSIS) OR XIAP OR HIAP-1 OR HILP OR (X-LINKED IAP) OR MIHA
S2	257315	ANTISENSE OR RIBOZYME OR TRIPLEX OR SIRNA OR RNAI OR (SHORT INTERFER? RNA) OR (RNA INTERFER?)
S3	10136532	CANCER? OR TUMOR? OR TUMOUR? OR CARCINOMA?
S4	1886463	INFLAMMATORY OR INFLAMMATION
S5	1147	(FOLLICULAR ATRESIA?) OR (ATRESIA? FOLLICULAR)
S6	515	S1 AND S2
S7	436	S6 AND S3
S8	3487542	6 AND S3 OR S4 OR S5
S9	1888026	S6 AND S3 OR S4 OR S5
S10	26	S6 AND S4
S11	2	S6 AND S5
S12	0	9 RD
S13	3487542	6 AND S3 OR S4 OR S5
S14	1888026	S6 AND S3 OR S4 OR S5
S15	436	S6 AND S3
S16	165	RD (unique items)
S17	10	S16 NOT PY>2000
S18	6121	(X-LINKED INHIBITOR OF APOPTOSIS) OR XIAP OR HIAP-1 OR HILP OR (X-LINKED IAP) OR MIHA
S19	257315	ANTISENSE OR RIBOZYME OR TRIPLEX OR SIRNA OR RNAI OR (SHORT INTERFER? RNA) OR (RNA INTERFER?)
S20	10136532	CANCER? OR TUMOR? OR TUMOUR? OR CARCINOMA?
S21	1886463	INFLAMMATORY OR INFLAMMATION
S22	1147	(FOLLICULAR ATRESIA?) OR (ATRESIA? FOLLICULAR)
S23	515	S18 AND S19
S24	436	S23 AND S20
S25	3487542	6 AND S20 OR S21 OR S22
S26	1888026	S23 AND S20 OR S21 OR S22
S27	26	S23 AND S21
S28	2	S23 AND S22
S29	0	9 RD
S30	3487542	6 AND S20 OR S21 OR S22
S31	1888026	S23 AND S20 OR S21 OR S22
S32	436	S23 AND S20
S33	165	RD (unique items)
S34	10	S33 NOT PY>2000

? T S27/MEDIUM,K/ALL

>>>KWIC option is not available in file(s): 398, 399

27/K/1 (Item 1 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.

0013914248 BIOSIS NO.: 200200507759

CD40 engagement enhances eosinophil survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

AUTHOR: Bureau Fabrice (Reprint); Seumois Gregory; Jaspar Fabrice;
Vanderplasschen Alain; Detry Bruno; Pastoret Paul-Pierre; Louis Renaud;
Lekeux Pierre

AUTHOR ADDRESS: Department of Physiology, Faculty of Veterinary Medicine,
University of Liege, Sart Tilman, Bat. B42, B-4000, Liege, Belgium**
Belgium

JOURNAL: Journal of Allergy and Clinical Immunology 110 (3): p443-449
September, 2002 2002

MEDIUM: print

ISSN: 0091-6749

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

...survival through induction of cellular inhibitor of apoptosis protein 2
expression: Possible involvement in allergic inflammation

...ABSTRACT: of the inhibitor of apoptosis protein (IAP) family, namely
cellular (c)-IAP1, c-IAP2, and **XIAP**, and 2 antiapoptotic proteins of
the Bcl-2 family, namely Bcl-xL and Bfl-1...

...staining with propidium iodide and FITC-conjugated annexin-V. c-IAP2
expression was inhibited with **antisense** oligonucleotides. Results:
Freshly isolated eosinophils from healthy and asthmatic patients did not
express CD40. Conversely...

...Inhibition of eosinophil apoptosis was accompanied by induction of
c-IAP2 but not c-IAP1, **XIAP**, Bcl-xL, or Bfl-1/Al expression. **Antisense**
knockdown of c-iap2 abolished CD40-induced enhancement of eosinophil
survival. Sputum cells from asthmatic...

...through induction of c-IAP2 expression and suggest a role for this
mechanism in allergic **inflammation**.

DESCRIPTORS:

DISEASES: allergic **inflammation** --

CHEMICALS & BIOCHEMICALS: ...X-inhibitor of apoptosis protein { **XIAP** };

27/K/2 (Item 1 from file: 440)
DIALOG(R)File 440:Current Contents Search(R)
(c) 2004 Inst for Sci Info. All rts. reserv.

19160771 Document Delivery Available: 0002235546

ISSN: 0021-9258

JOURNAL: JOURNAL OF BIOLOGICAL CHEMISTRY, 2004

(TABLE OF CONTENTS RECORD)

(The Complete Table of Contents now Available in Format 19)

...kappa B kinase (IKK) inhibitor, NEMO-binding domain peptide, blocks osteoclastogenesis and bone erosion in **inflammatory** arthritis. Dai S; Hirayama T; Abbas S; Abu-Amer Y. Washington Univ, Dept Orthoped Surg...

...000223554600075

- P. 37982-37996. Post-transcriptional regulation of endothelial nitric-oxide synthase by an overlapping **antisense** mRNA transcript. Robb GB; Carson AR; Tai SC; Fish JE; Singh S; Yamada T; Scherer...Geneva//Switzerland/. English. ARTICLE. 77 REFERENCES. ABSTRACT AVAILABLE. Document Delivery no: 000223554600028
- P. 37431-37435. **RNAi** -based analysis of CAP, Cbl, and CrkII function in the regulation of GLUT4 by insulin...processing of HtrA2/Omi is essential for induction of caspase-dependent cell death through antagonizing **XIAP**. Seong YM; Choi JY; Park HJ; Kim KJ; Ahn SG; Seong GH; Kim IK; Kang...

27/K/3 (Item 2 from file: 440)

DIALOG(R)File 440:Current Contents Search(R)
(c) 2004 Inst for Sci Info. All rts. reserv.

18293256 Document Delivery Available: 0002208704
ISSN: 0021-9258

JOURNAL: JOURNAL OF BIOLOGICAL CHEMISTRY , 2004
(TABLE OF CONTENTS RECORD)

(The Complete Table of Contents now Available in Format 19)

...AVAILABLE. Document Delivery no: 000220870400117

- P. 18091-18097. In vivo chromatin remodeling events leading to **inflammatory** gene transcription under diabetic conditions. Miao F; Gonzalo IG; Lanting L; Natarajan R. City Hope...

...Determination of the role of the human RNase H1 in the pharmacology of DNA-like **antisense** drugs. Wu HJ; Lima WF; Zhang H; Fan A; Sun H; Crooke ST. ISIS Pharmaceut...DIABLO selectively reduces the levels of c-IAP1 and c-IAP2 but not that of **XIAP** and livin in HeLa cells. Yang QH; Du CY. Stowers Inst Med Res, 1000E 50th...

27/K/4 (Item 3 from file: 440)

DIALOG(R)File 440:Current Contents Search(R)
(c) 2004 Inst for Sci Info. All rts. reserv.

16540078 Document Delivery Available: 0001841087
ISSN: 1078-0432

JOURNAL: CLINICAL CANCER RESEARCH , 2003
(TABLE OF CONTENTS RECORD)

(The Complete Table of Contents now Available in Format 19)

...English. ARTICLE. 25 REFERENCES. ABSTRACT AVAILABLE. Document Delivery no: 000184108700017

- P. 2510-2519. Efficacy of **antisense** morpholino oligomer targeted to c-myc in prostate cancer xenograft murine model and a phase...
AVAILABLE. Document Delivery no: 000184108700052

P. 2798-2806. The novel synthetic triterpenoid, CDDO-imidazolidine, inhibits **inflammatory** response and tumor growth in vivo. Place AE; Suh N; Williams CR; Risingsong R; Honda...Bloomington//MN/. English. ARTICLE. 46 REFERENCES. ABSTRACT AVAILABLE. Document Delivery no: 000184108700055

P. 2826-2836. **Antisense** oligonucleotides targeting **XIAP** induce apoptosis and enhance chemotherapeutic activity against human lung cancer cells in vitro and in...

...Utrecht//Netherlands/. English. ARTICLE. 40 REFERENCES. ABSTRACT AVAILABLE. Document Delivery no: 000184108700058

P. 2856-2865. **Antisense** Bcl-xl down-regulation switches the response to topoisomerase I inhibition from senescence to apoptosis...

27/K/5 (Item 4 from file: 440)

DIALOG(R) File 440:Current Contents Search(R)

(c) 2004 Inst for Sci Info. All rts. reserv.

14656002 Document Delivery Available: 000177936900015 References: 31

TITLE: CD40 engagement enhances eosinophil survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

AUTHOR(S): Bureau F (REPRINT); Seumois G; Jaspar F; Vanderplasschen A; Detry B; Pastoret PP; Louis R; Lekeux P

CORPORATE SOURCE: Univ Liege, Dept Physiol, Bat B42/B-4000 Liege//Belgium/ (REPRINT); Univ Liege, Dept Physiol, /B-4000 Liege//Belgium/; Univ Liege, Dept Immunol Vaccinol, /B-4000 Liege//Belgium/; Univ Liege, Dept Pneumol, /B-4000 Liege//Belgium/

PUBLICATION TYPE: JOURNAL

PUBLICATION: JOURNAL OF ALLERGY AND CLINICAL IMMUNOLOGY, 2002, V110, N3 (SEP), P443-449

GENUINE ARTICLE#: 592KM

PUBLISHER: MOSBY, INC, 11830 WESTLINE INDUSTRIAL DR, ST LOUIS, MO 63146-3318 USA

ISSN: 0091-6749

LANGUAGE: English **DOCUMENT TYPE:** ARTICLE (ABSTRACT AVAILABLE)

...**TITLE: survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation**

...**ABSTRACT:** of the inhibitor of apoptosis protein (IAP) family, namely cellular (c)-IAP1, c-IAP2, and **XIAP**, and 2 antiapoptotic proteins of the Bcl-2 family, namely Bcl-x(L) and Bfl...

...staining with propidium iodide and FITC-conjugated annexin-V. c-IAP2 expression was inhibited with **antisense** oligonucleotides.

Results: Freshly isolated eosinophils from healthy and asthmatic patients did not express CD40. Conversely...

...Inhibition of eosinophil apoptosis was accompanied by induction of c-IAP2 but not c-IAP1, **XIAP**, Bcl-x(L), or Bfl-1/A1 expression. **Antisense** knockdown of c-iap2 abolished CD40-induced enhancement of eosinophil survival. Sputum cells from asthmatic...

...through induction of c-IAP2 expression and suggest a role for this mechanism in allergic inflammation .

27/K/6 (Item 5 from file: 440)

DIALOG(R)File 440:Current Contents Search(R)
(c) 2004 Inst for Sci Info. All rts. reserv.

12111850

ISSN: 0021-9258

JOURNAL: JOURNAL OF BIOLOGICAL CHEMISTRY , 2000

(TABLE OF CONTENTS RECORD)

(The Complete Table of Contents now Available in Format 19)

...Div, /E Melbourne/Vic 3000/Australia/. English. ARTICLE. 33
REFERENCES. ABSTRACT AVAILABLE

P. 32077-32088. **Inflammatory** versus proliferative processes in epidermis - Tumor necrosis factor alpha induces K6b keratin synthesis through a...Dept Biomed Engn, /Memphis//TN/38152. English. ARTICLE. 50 REFERENCES. ABSTRACT AVAILABLE

P. 31733-31738. **XIAP** regulates DNA damage-induced apoptosis downstream of caspase-9 cleavage. Datta R; ...D-35033 Marburg//Germany/. English. ARTICLE. 58 REFERENCES. ABSTRACT AVAILABLE

P. 32281-32288. Expression of **antisense** to integrin subunit beta(3) inhibits microvascular endothelial cell capillary tube formation in fibrin. Dallabrida...

27/K/7 (Item 1 from file: 34)

DIALOG(R)File 34:SciSearch(R) Cited Ref Sci
(c) 2004 Inst for Sci Info. All rts. reserv.

10980082 Genuine Article#: 592KM No. References: 31

Title: CD40 engagement enhances eosinophil survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

Author(s): Bureau F (REPRINT) ; Seumois G; Jaspar F; Vanderplasschen A; Detry B; Pastoret PP; Louis R; Lekeux P

Corporate Source: Univ Liege,Fac Vet Med, Dept Physiol,Bat B42/B-4000 Liege//Belgium/ (REPRINT); Univ Liege,Fac Vet Med, Dept Physiol,B-4000 Liege//Belgium/; Univ Liege,Fac Vet Med, Dept Immunol Vaccinol,B-4000 Liege//Belgium/; Univ Liege,Fac Med, Dept Pneumol,B-4000 Liege//Belgium/

Journal: JOURNAL OF ALLERGY AND CLINICAL IMMUNOLOGY, 2002, V110, N3 (SEP), P443-449

ISSN: 0091-6749 Publication date: 20020900

Publisher: MOSBY, INC, 11830 WESTLINE INDUSTRIAL DR, ST LOUIS, MO 63146-3318 USA

Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE)

...Title: survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

...Abstract: of the inhibitor of apoptosis protein (IAP) family, namely

cellular (c)-IAP1, c-IAP2, and **XIAP** , and 2 antiapoptotic proteins of the Bcl-2 family, namely Bcl-x(L) and Bfl...

...staining with propidium iodide and FITC-conjugated annexin-V. c-IAP2 expression was inhibited with **antisense** oligonucleotides.

Results: Freshly isolated eosinophils from healthy and asthmatic patients did not express CD40. Conversely...

...Inhibition of eosinophil apoptosis was accompanied by induction of c-IAP2 but not c-IAP1, **XIAP** , Bcl-x(L), or Bfl-1/A1 expression.

Antisense knockdown of c-iap2 abolished CD40-induced enhancement of eosinophil survival. Sputum cells from asthmatic...

...through induction of c-IAP2 expression and suggest a role for this mechanism in allergic **inflammation** .

27/K/8 (Item 1 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

11994848 PMID: 12209092

CD40 engagement enhances eosinophil survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation .

Bureau Fabrice; Seumois Gregory; Jaspar Fabrice; Vanderplasschen Alain; Detry Bruno; Pastoret Paul-Pierre; Louis Renaud; Lekeux Pierre
Department of Physiology, Faculty of Veterinary Medicine, University of Liege, Belgium.

Journal of allergy and clinical immunology (United States) Sep 2002, 110 (3) p443-9, ISSN 0091-6749 Journal Code: 1275002

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

...survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation .

... of the inhibitor of apoptosis protein (IAP) family, namely cellular (c)-IAP1, c-IAP2, and **XIAP** , and 2 antiapoptotic proteins of the Bcl-2 family, namely Bcl-x(L) and Bfl...

... staining with propidium iodide and FITC-conjugated annexin-V. c-IAP2 expression was inhibited with **antisense** oligonucleotides. RESULTS: Freshly isolated eosinophils from healthy and asthmatic patients did not express CD40. Conversely...

... Inhibition of eosinophil apoptosis was accompanied by induction of c-IAP2 but not c-IAP1, **XIAP** , Bcl-x(L), or Bfl-1/A1 expression. **Antisense** knockdown of c-iap2 abolished CD40-induced enhancement of eosinophil survival. Sputum cells from asthmatic...

... through induction of c-IAP2 expression and suggest a role for this mechanism in allergic **inflammation** .

...; IM; Cell Survival; Cells, Cultured; Eosinophilia--immunology--IM; Eosinophils--cytology--CY; Hypersensitivity, Immediate--pathology--PA; **Inflammation** --immunology--IM; Sputum--cytology--CY

27/K/9 (Item 1 from file: 399)
DIALOG(R)File 399:CA SEARCH(R)
(c) 2004 American Chemical Society. All rts. reserv.

141307497 CA: 141(19)307497m PATENT

Use of caspase inhibitors as antiviral agents, and test system for their discovery

INVENTOR(AUTHOR): Ludwig, Stefan; Planz, Oliver; Sedlacek, Hans-Harald; Pleschka, Stephan

LOCATION: Germany,

ASSIGNEE: Medinnova Gesellschaft fur Medizinische Innovationen aus Akademischer Forschung m.b.H.

PATENT: PCT International ; WO 200485682 A2 DATE: 20041007

APPLICATION: WO 2004DE646 (20040324) *DE 10313636 (20030326)

PAGES: 40 pp. CODEN: PIXXD2 LANGUAGE: German CLASS: C12Q-001/70A

DESIGNATED COUNTRIES: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG; BR; BW; BY; BZ; CA; CH; CN; CO; CR; CU; CZ; DE; DK; DM; DZ; EC; EE; EG; ES; FI; GB; GD; GE; GH; GM; HR; HU; ID; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR; LS; LT; LU; LV; MA; MD; MG; MK; MN; MW; MX; MZ; NA; NI; NO; NZ; OM; PG; PH; PL; PT; RO; RU; SC; SD; SE; SG; SK; SL; SY; TJ; TM; TN; TR; TT; TZ; UA; UG; US; UZ; VC; VN; YU; ZA; ZM; ZW DESIGNATED REGIONAL: BW; GH; GM; KE; LS; MW; MZ; SD; SL; SZ; TZ; UG; ZM; ZW; AM; AZ; BY; KG; KZ; MD; RU; TJ; TM; AT; BE; BG; CH; CY; CZ; DE; DK; EE; ES; FI; FR; GB; GR; HU; IE; IT; LU; MC; NL; PL; PT; RO; SE; SI; SK; TR; BF; BJ; CF; CG; CI; CM; GA; GN; GQ; GW; ML; MR; NE; SN; TD; TG

27/K/10 (Item 1 from file: 73)
DIALOG(R)File 73:EMBASE
(c) 2004 Elsevier Science B.V. All rts. reserv.

11770551 EMBASE No: 2002339725

CD40 engagement enhances eosinophil survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

Bureau F.; Seumois G.; Jaspar F.; Vanderplasschen A.; Detry B.; Pastoret P.-P.; Louis R.; Lekeux P.

Dr. F. Bureau, Department of Physiology, Faculty of Veterinary Medicine, University of Liege, Sart Tilman, B-4000 Liege Belgium
Journal of Allergy and Clinical Immunology (J. ALLERGY CLIN. IMMUNOL.)
(United States) 2002, 110/3 (443-449)

CODEN: JACIB ISSN: 0091-6749

DOCUMENT TYPE: Journal ; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 31

...survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

...of the inhibitor of apoptosis protein (IAP) family, namely cellular (c)-IAP1, c-IAP2, and XIAP, and 2 antiapoptotic proteins of the Bcl-2 family, namely Bcl-xSUBL and Bfl-1...

...staining with propidium iodide and FITC-conjugated annexin-V, c-IAP2 expression was inhibited with antisense oligonucleotides. Results: Freshly isolated eosinophils from healthy and asthmatic patients did not express CD40. Conversely...

...Inhibition of eosinophil apoptosis was accompanied by induction of c-IAP2 but not c-IAP1, **XIAP**, Bcl-xSUBL, or Bfl-1/A1 expression.

Antisense knockdown of c-iap2 abolished CD40-induced enhancement of eosinophil survival. Sputum cells from asthmatic...

...through induction of c-IAP2 expression and suggest a role for this mechanism in allergic **inflammation**.

DRUG DESCRIPTORS:

propidium iodide; fluorescein isothiocyanate; lipocortin 5; **antisense** oligonucleotide; RNA--endogenous compound--ec; unclassified drug

MEDICAL DESCRIPTORS:

*allergic asthma--etiology--et; * **inflammation** --etiology--et; *eosinophil

27/K/11 (Item 1 from file: 71)

DIALOG(R)File 71:ELSEVIER BIOBASE

(c) 2004 Elsevier Science B.V. All rts. reserv.

02138183 2002219133

CD40 engagement enhances eosinophil survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

Bureau F.; Seumois G.; Jaspar F.; Vanderplasschen A.; Detry B.; Pastoret P.-P.; Louis R.; Lekeux P.

ADDRESS: Dr. F. Bureau, Department of Physiology, Faculty of Veterinary Medicine, University of Liege, Sart Tilman, B-4000 Liege, Belgium

Journal: Journal of Allergy and Clinical Immunology, 110/3 (443-449), 2002, United States

CODEN: JACIB

ISSN: 0091-6749

DOCUMENT TYPE: Article

LANGUAGES: English SUMMARY LANGUAGES: English

NO. OF REFERENCES: 31

DESCRIPTORS:

Allergy; Apoptosis; Asthma; Atopy; CD40; Eosinophils; Granulocytes; **Inflammation**; Neutrophils

CLASSIFICATION CODE AND DESCRIPTION:

86.3.3.9 - IMMUNOLOGY AND INFECTIOUS DISEASES / CELLS OF THE IMMUNE SYSTEM / T Lymphocytes / General functions and activation

86.3.9 - IMMUNOLOGY AND INFECTIOUS DISEASES / CELLS OF THE IMMUNE SYSTEM / Eosinophil Leucocytes

86.8.1.4 - IMMUNOLOGY AND INFECTIOUS DISEASES / IMMUNE RESPONSE DISORDERS / Acute **Inflammation**, Immediate Hypersensitivity, Anaphylaxis / Diagnosis and therapy

89.2.5.4 - CELL AND DEVELOPMENTAL BIOLOGY / CELL GROWTH AND DIVISION / Cellular Senescence and Death / Death (apoptosis)

...survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

...of the inhibitor of apoptosis protein (IAP) family, namely cellular (c)-IAP1, c-IAP2, and **XIAP**, and 2 antiapoptotic proteins of the Bcl-2 family, namely Bcl-xSUBL and Bfl-1...

...staining with propidium iodide and FITC-conjugated annexin-V, c-IAP2 expression was inhibited with **antisense** oligonucleotides. Results:

Freshly isolated eosinophils from healthy and asthmatic patients did not express CD40. Conversely...

...Inhibition of eosinophil apoptosis was accompanied by induction of c-IAP2 but not c-IAP1, **XIAP**, Bcl-xSUBL, or Bfl-1/A1 expression.

Antisense knockdown of c-iap2 abolished CD40-induced enhancement of eosinophil survival. Sputum cells from asthmatic...

...through induction of c-IAP2 expression and suggest a role for this mechanism in allergic **inflammation**.

DESCRIPTORS:

Allergy; Apoptosis; Asthma; Atopy; CD40; Eosinophils; Granulocytes; **Inflammation**; Neutrophils

CLASSIFICATION CODE AND DESCRIPTION:

...Acute **Inflammation**, Immediate Hypersensitivity, Anaphylaxis...

27/K/12 (Item 1 from file: 144)

DIALOG(R) File 144:Pascal

(c) 2004 INIST/CNRS. All rts. reserv.

15809300 PASCAL No.: 02-0525803

CD40 engagement enhances eosinophil survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

BUREAU Fabrice; SEUMOIS Gregory; JASPAR Fabrice; VANDERPLASSCHEN Alain; DETRY Bruno; PASTORET Paul-Pierre; LOUIS Renaud; LEKEUX Pierre

Department of Physiology, Faculty of Veterinary Medicine, Department of Pneumology, Faculty of Medicine, University of Liege, Liege, Belgium; Department of Immunology/Vaccinology, Faculty of Veterinary Medicine, Department of Pneumology, Faculty of Medicine, University of Liege, Liege, Belgium

Journal: Journal of allergy and clinical immunology, 2002, 110 (3) 443-449

Language: English

Copyright (c) 2002 INIST-CNRS. All rights reserved.

...survival through induction of cellular inhibitor of apoptosis protein 2 expression: Possible involvement in allergic inflammation

... of the inhibitor of apoptosis protein (IAP) family, namely cellular (c)-IAP1, c-IAP2, and **XIAP**, and 2 antiapoptotic proteins of the Bcl-2 family, namely Bcl-x SUB L and...

... staining with propidium iodide and FITC-conjugated annexin-V. c-IAP2 expression was inhibited with **antisense** oligonucleotides. Results: Freshly isolated eosinophils from healthy and asthmatic patients did not express CD40. Conversely...

... Inhibition of eosinophil apoptosis was accompanied by induction of c-IAP2 but not c-IAP1, **XIAP**, Bcl-x SUB L, or Bfl-1/A1 expression.

Antisense knockdown of c-iap2 abolished CD40-induced enhancement of eosinophil survival. Sputum cells from asthmatic...

... through induction of c-IAP2 expression and suggest a role for this mechanism in allergic **inflammation**.

English Descriptors: Asthma; Allergy; Atopy; Human; Pathogenesis;
Eosinophil; **Inflammation** ; Granulocyte; Apoptosis; Blood; Sputum; Flow
cytometry; Immunoblotting assay

French Descriptors: Asthme; Allergie; Atopie; Homme; Pathogenie;
Eosinophile; **Inflammation** ; Granulocyte; Apoptose; Sang; Expectoration;
Cytometrie flux; Methode immunoblotting; Antigene CD40; Proteine IAP

27/K/13 (Item 2 from file: 144)
DIALOG(R) File 144:Pascal
(c) 2004 INIST/CNRS. All rts. reserv.

14624714 PASCAL No.: 00-0295143
**Gene therapy that inhibits nuclear translocation of nuclear factor KB
results in tumor necrosis factor alpha -induced apoptosis of human
synovial fibroblasts**
ZHANG H G; NING HUANG; DI LIU; BILBAO L; XIAOWU ZHANG; YANG P; TONG ZHOU;
CURIEL D T; MOUNTZ J D
University of Alabama at Birmingham, United States; Gene Therapy Program,
Birmingham, Alabama, United States; University of Alabama at Birmingham,
Birmingham, Alabama, United States; Veterans Administration Medical Center,
Birmingham, Alabama, United States
Journal: Arthritis and rheumatism, 2000, 43 (5) 1094-1105
Language: English

Copyright (c) 2000 INIST-CNRS. All rights reserved.

... factor KB (IKB) dominant-negative adenovirus (AdCMVIKB-DN) and an
X-linked inhibitor of apoptosis (**XIAP**) **antisense** adenovirus
(AdCMVXIAP-AS). Primary RA synovial fibroblast (RASf) cell lines were
transfected in vitro, and...

... was no apoptosis after treatment with AdCMVI kappa B-DN in the absence
of TNFa. **XIAP** is an inhibitor of apoptosis which was up-regulated by
TNFa, and this up-regulation...

... greatly enhances apoptosis due to inhibition of an NF-KB-mediated
antiapoptosis signaling pathway, and **XIAP** is a TNFa-inducible specific
inhibitor of apoptosis in RA synovial cell lines. This and...

Broad Descriptors: Diseases of the osteoarticular system; **Inflammatory**
joint disease; Immunopathology; Autoimmune disease; Cytokine; Systeme
osteoarticulaire pathologie; Rhumatisme inflammatoire; Immunopathologie;
Maladie autoimmune; Cytokine...

27/K/14 (Item 1 from file: 991)
DIALOG(R) File 991:NewsRoom 2004 Jan 1-2004/Jul 31
(c) 2004 The Dialog Corporation. All rts. reserv.

0836083099 16L82K4U
POSTERS: SIGNAL TRANSDUCTION (NOT INSULIN ACTION)-CYTOKINES AND APOPTOSIS
Cao, Wen M
Murao, Koji
Imachi, Hitomi
Yu, Xiao
Et al
Diabetes, pA442

Tuesday, June 1, 2004

JOURNAL CODE: AHKF LANGUAGE: English RECORD TYPE: Fulltext

DOCUMENT TYPE: Trade Journal ISSN: 0012-1797

WORD COUNT: 3,312

...protein bound the PRCE by EMSA and was identified by super shift assay. Finally, PREB- **siRNA** , that inhibited the PREB expression, significantly decreased the effect of glucose on MCP-1 promoter...

...FN mRNA and protein expression in a dose-dependent manner. HUVECs were then transfected with **SiRNA** targeted to oncofetal FN. The transfected endothelial cells were cultured in low (5 mmol/L)...
...glucose caused upregulation of VEGF mRNA levels in the negative control transfected cells. Transfection with **SiRNA** targeted to oncofetal FN completely abolished glucose-induced VEGF mRNA expression. Endothelial cells were then...

...K. KIM. New Haven, CT

We have previously shown that acute pre-treatment of anti- **inflammatory** cytokine, interleukin (IL)-10, prevented insulin resistance associated with 5-hour lipid infusion, and the...

...Research

1859-P

Cytoprotection of Pancreatic [beta] Cells by Co-Overexpression of Bcl-2 and **XIAP** Genes Blocking TRAIL-Mediated Death-Signal Transduction Pathways

DAWEI OU, XIAJIE WANG, DANIEL L. METZGER...

...investigated the inhibitory effects of Bcl-2 and the X-linked inhibitor of apoptosis protein (**XIAP**) on TNF-related apoptosis-inducing ligand (TRAIL)-induced human [beta]-cell destruction. A panel of...

...type NES2Y and CM cells and the vector transfectants (P<0.0002 to 0.01). **XIAP** -overexpressed NES2Y and CM cells were developed by exposing the cells to an infectious **XIAP** recombinant adenovirus (AdXIAP) at MOI 10. [beta] cells infected with AdLacZ were used as controls...

...AdXIAP were much less than that observed in cells transfected with either Bcl-2 or **XIAP** alone (P<0.0004 to 0.04). Overexpression of Bcl-2 and/or **XIAP** by gene transfection inhibited TRAIL-induced activation of caspases and TRAIL-mediated damage of mitochondrial...

...these cells suggesting the major mechanisms of regulation. These results indicated that Bcl-2 and **XIAP** are important components of the signal transduction pathways that regulate TRAIL-induced human [beta]-cell death and that there may be novel therapeutic potential of Bcl-2 and **XIAP** co-overexpression [beta] cells in type 1 diabetes.

1860-P

Protein Kinase C alpha Activation...

DIALOG(R)File 991:NewsRoom 2004 Jan 1-2004/Jul 31
(c) 2004 The Dialog Corporation. All rts. reserv.

0803005993 16J605V8

OTHER NEWS TO NOTE

BioWorld Today, v15

Tuesday, March 30, 2004

JOURNAL CODE: ARME LANGUAGE: English RECORD TYPE: Fulltext

DOCUMENT TYPE: Newsletter SECTION HEADING: 60 ISSN: 1541 0595

WORD COUNT: 1,824

Aegera Therapeutics Inc., of Montreal, began Phase I trials of AEG35156/GEM640, an **antisense** inhibitor of x-linked inhibitor of apoptosis protein (**XIAP**), in collaboration with Cancer Research UK. AEG35156 is a second-generation **antisense** oligonucleotide, designed to reduce high levels of **XIAP** in cancer cells and enable cell death. The initial trial is in patients with advanced...

...agreement with Funakoshi Ltd., of Tokyo, related to RNAX's services for the validation of **siRNA** oligonucleotides and genetic targets based on RNA interference technology. Funakoshi will act as exclusive agent...

...applications for endothelial differentiation gene (Edg) receptors 1 through 7 and their use in cancer, **inflammation** and other diseases. Ceretek also transferred its cell lines incorporating chimeric Edg receptors for screening...

27/K/16 (Item 1 from file: 992)

DIALOG(R)File 992:NewsRoom 2003

(c) 2004 The Dialog Corporation. All rts. reserv.

0589510171 162V09XU

Research and Markets: Targeting endogenous inhibitors of apoptosis: Opportunities for the treatment of cancer, stroke and MS

M2 Communications

Tuesday, February 11, 2003

JOURNAL CODE: ALPP LANGUAGE: English RECORD TYPE: Fulltext

DOCUMENT TYPE: Newswire

WORD COUNT: 1,549

...Proteins") family. This field has grown exponentially since 1995 and continues to do so. Although **XIAP** and survivin remain the better known members of this family, 8 human IAPs have now...

...contributed much of the data surrounding one of the most promising targets from this family, **XIAP** .

Dr Jon Goldhill: Dr Jon Goldhill has over 10 years of academic and industrial research...

...management at the French pharmaceutical giants, Sanofi-Synthelabo. Focussing on a variety of indications including **inflammatory** disorders, GI disease, Urological conditions and cancer, Dr Goldhill was responsible for target identification and....

...apoptosis inhibitory protein; NAIP) BIRC2 (API1; HIAP2; cIAP1; MIHB) BIRC3 (API2; HIAP1; cIAP2; MIHC) BIRC4 (**XIAP** ; API3; **MIHA** ; ILP) BIRC5

(Survivin; API4; TLAP) BIRC6 (Apollon; BRUCE) BIRC7 (MLIAP; KIAP; Livin)
BIRC8: (ILP-2...

...13-PE38 apolizumab alvocidib indisulam combretastatin A-4 Profiles of
IAP-related molecules HIAP-1 **antisense XIAP** inhibitors AEG-161
Apoptosis inhibitors Trends in apoptosis inhibitors Apoptosis inhibitors in
development Profiles of...

27/K/17 (Item 1 from file: 20)
DIALOG(R)File 20:Dialog Global Reporter
(c) 2004 The Dialog Corp. All rts. reserv.

27495364 (USE FORMAT 7 OR 9 FOR FULLTEXT)
**Research and Markets: Targeting endogenous inhibitors of apoptosis:
Opportunities for the treatment of cancer, stroke and MS**
M2 PRESSWIRE
February 11, 2003
JOURNAL CODE: WMPR LANGUAGE: English RECORD TYPE: FULLTEXT
WORD COUNT: 1471

(USE FORMAT 7 OR 9 FOR FULLTEXT)

... Proteins") family. This field has grown exponentially since 1995
and continues to do so. Although **XIAP** and survivin remain the better
known members of this family, 8 human IAPs have now...

... contributed much of the data surrounding one of the most promising
targets from this family, **XIAP** .

Dr Jon Goldhill: Dr Jon Goldhill has over 10 years of academic and
industrial research...

... management at the French pharmaceutical giants, Sanofi-Synthelabo.
Focussing on a variety of indications including **inflammatory** disorders,
GI disease, Urological conditions and cancer, Dr Goldhill was responsible
for target identification and...

... apoptosis inhibitory protein; NAIP) BIRC2 (API1; HIAP2; cIAP1; MIHB)
BIRC3 (API2; HIAP1; cIAP2; MIHC) BIRC4 (**XIAP** ; API3; **MIHA** ; ILP) BIRC5
(Survivin; API4; TLAP) BIRC6 (Apollon; BRUCE) BIRC7 (MLIAP; KIAP; Livin)
BIRC8: (ILP-2...

... 13-PE38 apolizumab alvocidib indisulam combretastatin A-4 Profiles of
IAP-related molecules HIAP-1 **antisense XIAP** inhibitors AEG-161
Apoptosis inhibitors Trends in apoptosis inhibitors Apoptosis inhibitors in
development Profiles of...

27/K/18 (Item 1 from file: 761)
DIALOG(R)File 761:Datamonitor Market Res.
(c) 2004 Datamonitor. All rts. reserv.

00241838

CANCER: 1.1 THIS MONTH'S HIGHLIGHTS

Main Title: PHARMAWATCH
Pub. Date: July 23, 2003
Source: DATAMONITOR

Telephone: +44 20 7675 7000
Word Count: 3689 (2 pp.)
Language: English

Features: TABLE
Country: WORLD
Industry: HEALTH CARE

Company Names (DIALOG Generated): Accentia Inc ; American Association for Cancer Research ; American Cancer Society ; American Gastroenterological Association ; American Society of Clinical Oncology ; American Society of Hematology ; American Urological Association ; Amgen ; AnorMed ; Ariad Pharmaceuticals ; AstraZeneca ; Aton Pharma ; ARIA ; AVI BioPharma ; Bayer ; Beatson Cancer Institute ; Beatson Oncology Centre ; Biogen Idec Inc ; Biologics License Application ; Biomira ; Biovest International ; BioTransplant ; Bone Metastases ; Breast Cancer Center of Excellence ; Bristol Myers Squibb ; British Biotech ; BMS/Celltech ; BMJ ; Cancer Treatment ; Case Western Reserve University School of Medicine ; Catholic University ; Cell Therapeutics ; Celltech ; Chugai ; Committee for Proprietary Medicinal ; Corixa Corporation ; Department of Defense ; Drug Safety Monitoring Committee ; DNA ; Eli Lilly and Company ; Equal ; European Agency for the Evaluation of Medicinal Products ; European Commission ; European Committee on Proprietary Medicinal ; European Neurological Society ; Genentech ; Genta ; Georgetown University Medical Center ; German Federal Institute for Drugs ; GlaxoSmithKline ; GASTRO ; GTx Inc ; GTAC ; GTI ; Idec Biogen ; Idec Pharmaceuticals ; Immunex ; Immunology ; ImClone Systems ; Indre et Loire ; Investigational New Drug ; Ireland Cancer Center ; Ivax Corporation ; IVX ; Journal ; Leicester Royal Infirmary ; Leukemia & Lymphoma Society ; Lombardi Cancer Center ; Marketing Authorisation Application ; Medical Devices ; Medicinal Products ; MedImmune ; Memorial Sloan Kettering Cancer Center ; Merck & Co ; Merck KGaA ; Merger & Acquisition Update ; Millennium Pharmaceutical ; MAA ; MD Anderson Cancer Center ; ML ; National Cancer Institute ; Nemod Immunotherapie AG ; Netherlands Cancer Institute ; New Drug Application ; New England Journal ; Novuspharma SpA ; Nurses 's Health ; Ohio State University Medical Center ; Orphan Drug ; Other Cancers News ; Oxford BioMedica ; Pfizer ; PharmaMar ; Point ; Praecis Pharmaceuticals ; Progen Industries ; PGLAF ; PSA ; Regulatory News ; Reuters ; Roche ; Royal Liverpool University Hospital ; Royal Marsden Hospital ; Sanofi Synthelabo Research ; Southern Research Institute ; Stanford Research Institute International ; Stanford University ; State University of New York at Buffalo ; Sundsvall Hospital ; SuperGen ; Sustained Without M & A ; SRI International ; Teva Pharmaceutical ; Texas Health Sciences Center ; Texas MD Anderson Cancer Center ; University of Texas MD Anderson Cancer Center ; Titan Pharmaceuticals ; TransMolecular ; TAP ; University of Chicago ; University of Cologne Aventis Pharma and Mologen ; University of Texas Health Sciences Center ; University of Texas MD Anderson Cancer Center ; University of Wisconsin Medical School ; UK Gene Therapy Advisory Committee ; Washington Hospital Center ; Wisconsin Medical School ; Women 's Health Initiative

...must be provided, which will be caused by a non-specific immune activation similar to **inflammation** reactions. All three signals are

generated by dSLIM.

The study was of a five-armed...

...a tendency to interfere with similar enzymes called sheddases that play a part in secreting **inflammatory** signals. Sheddase inhibition can cause arthritis. BMS275291 was rationally designed using structural biology to aim...cancer target

Scientists from AVI BioPharma have presented research data describing the successful inhibition of **Xiap**, one of a family of genes known to inhibit apoptosis, or programmed cell death, in cancer cells using an **antisense** strategy.

The presentation highlighted results from a series of preclinical studies using AVI BioPharma's Neugene **antisense** technology to block the expression of the gene **Xiap**. In a wide variety of cancers, including prostate cancer, the production of **Xiap** helps tumors grow by preventing apoptosis, which would normally occur in cells damaged by chemotherapy...

...cells that have become resistant to chemotherapy typically exhibit anti-apoptotic behavior. By using Neugene **antisense** drugs to block the **Xiap** gene, the AVI scientists demonstrated that prostate cancer cells that previously became resistant to chemotherapy can be resensitized to chemotherapy drugs. The **Xiap** target was selected over other potential anti-apoptotic targets, such as BCL-2, because of...

...release. "This study builds on our previous experience in several cancer indications, using our Neugene **antisense** drugs to block select genes that are associated with cancer progression. Target selection is a critical component to **antisense** drug development for cancer, and with preclinical data for **Xiap** and androgen receptor targets and clinical data for the c-myc target, we believe we have assembled a robust **antisense** cancer program."

GTx's Acapodene shown to reduces high-grade prostate intraepithelial neoplasia
GTx, Inc...

...sponsored Investigational New Drug application for a clinical trial of Lorus Therapeutics' (LORFF.OB) lead **antisense** drug, GTI-2040 in combination with cytarabine, in patients with refractory or relapsed acute myeloid...three cancer therapies in Biogen's product pipeline. Having traditionally been focused on neurology and **inflammatory** diseases, Biogen looked to find a partner more experienced in oncology. It would seem that ...

27/K/19 (Item 2 from file: 761)

DIALOG(R)File 761:Datamonitor Market Res.

(c) 2004 Datamonitor. All rts. reserv.

00240994

CANCER: 1.1 THIS MONTH'S HIGHLIGHTS

Main Title: PHARMAWATCH

Pub. Date: July 23, 2003

Source: DATAMONITOR

Telephone: +44 20 7675 7000

Word Count: 3689 (2 pp.)

Language: English

Features: TABLE

Country: WORLD

Industry: HEALTH CARE

Company Names (DIALOG Generated): Accentia Inc ; American Association for Cancer Research ; American Cancer Society ; American Gastroenterological Association ; American Society of Clinical Oncology ; American Society of Hematology ; American Urological Association ; Amgen ; AnorMed ; Ariad Pharmaceuticals ; AstraZeneca ; Aton Pharma ; ARIA ; AVI BioPharma ; Bayer ; Beatson Cancer Institute ; Beatson Oncology Centre ; Biogen Idec Inc ; Biologics License Application ; Biomira ; Biovest International ; BioTransplant ; Bone Metastases ; Breast Cancer Center of Excellence ; Bristol Myers Squibb ; British Biotech ; BMS/Celltech ; BMY ; Cancer Treatment ; Case Western Reserve University School of Medicine ; Catholic University ; Cell Therapeutics ; Celltech ; Chugai ; Committee for Proprietary Medicinal ; Corixa Corporation ; Department of Defense ; Drug Safety Monitoring Committee ; DNA ; Eli Lilly and Company ; Equal ; European Agency for the Evaluation of Medicinal Products ; European Commission ; European Committee on Proprietary Medicinal ; European Neurological Society ; Genentech ; Genta ; Georgetown University Medical Center ; German Federal Institute for Drugs ; GlaxoSmithKline ; GASTRO ; GTx Inc ; GTAC ; GTI ; Idec Biogen ; Idec Pharmaceuticals ; Immunex ; Immunology ; ImClone Systems ; Indre et Loire ; Investigational New Drug ; Ireland Cancer Center ; Ivax Corporation ; IVX ; Journal ; Leicester Royal Infirmary ; Leukemia & Lymphoma Society ; Lombardi Cancer Center ; Marketing Authorisation Application ; Medical Devices ; Medicinal Products ; MedImmune ; Memorial Sloan Kettering Cancer Center ; Merck & Co ; Merck KgaA ; Merger & Acquisition Update ; Millennium Pharmaceutical ; MAA ; MD Anderson Cancer Center ; ML ; National Cancer Institute ; Nemod Immunotherapie AG ; Netherlands Cancer Institute ; New Drug Application ; New England Journal ; Novuspharma SpA ; Nurses 's Health ; Ohio State University Medical Center ; Orphan Drug ; Other Cancers News ; Oxford BioMedica ; Pfizer ; PharmaMar ; Point ; Praecis Pharmaceuticals ; Progen Industries ; PGLAF ; PSA ; Regulatory News ; Reuters ; Roche ; Royal Liverpool University Hospital ; Royal Marsden Hospital ; Sanofi Synthelabo Research ; Southern Research Institute ; Stanford Research Institute International ; Stanford University ; State University of New York at Buffalo ; Sundsvall Hospital ; SuperGen ; Sustained Without M & A ; SRI International ; Teva Pharmaceutical ; Texas Health Sciences Center ; Texas MD Anderson Cancer Center ; University of Texas MD Anderson Cancer Center ; Titan Pharmaceuticals ; TransMolecular ; TAP ; University of Chicago ; University of Cologne Aventis Pharma and Mologen ; University of Texas Health Sciences Center ; University of Texas MD Anderson Cancer Center ; University of Wisconsin Medical School ; UK Gene Therapy Advisory Committee ; Washington Hospital Center ; Wisconsin Medical School ; Women 's Health Initiative

...must be provided, which will be caused by a non-specific immune activation similar to **inflammation** reactions. All three signals are generated by dSLIM.

The study was of a five-armed...

...a tendency to interfere with similar enzymes called sheddases that play a part in secreting **inflammatory** signals. Sheddase inhibition can cause arthritis. BMS275291 was rationally designed using structural biology to aim...cancer target

Scientists from AVI BioPharma have presented research data describing the successful inhibition of **Xiap**, one of a family of genes known to inhibit apoptosis, or programmed cell death, in cancer cells using an **antisense** strategy.

The presentation highlighted results from a series of preclinical studies using AVI BioPharma's Neugene **antisense** technology to block the expression of the gene **Xiap**. In a wide variety of cancers, including prostate cancer, the production of **Xiap** helps tumors grow by preventing apoptosis, which would normally occur in cells damaged by chemotherapy...

...cells that have become resistant to chemotherapy typically exhibit anti-apoptotic behavior. By using Neugene **antisense** drugs to block the **Xiap** gene, the AVI scientists demonstrated that prostate cancer cells that previously became resistant to chemotherapy can be resensitized to chemotherapy drugs. The **Xiap** target was selected over other potential anti-apoptotic targets, such as BCL-2, because of...

...release. "This study builds on our previous experience in several cancer indications, using our Neugene **antisense** drugs to block select genes that are associated with cancer progression. Target selection is a critical component to **antisense** drug development for cancer, and with preclinical data for **Xiap** and androgen receptor targets and clinical data for the c-myc target, we believe we have assembled a robust **antisense** cancer program."

GTx's Acapodene shown to reduces high-grade prostate intraepithelial neoplasia
GTx, Inc...

...sponsored Investigational New Drug application for a clinical trial of Lorus Therapeutics' (LORFF.OB) lead **antisense** drug, GTI-2040 in combination with cytarabine, in patients with refractory or relapsed acute myeloid...three cancer therapies in Biogen's product pipeline. Having traditionally been focused on neurology and **inflammatory** diseases, Biogen looked to find a partner more experienced in oncology. It would seem that
...

27/K/20 (Item 3 from file: 761)

DIALOG(R)File 761:Datamonitor Market Res.
(c) 2004 Datamonitor. All rts. reserv.

00203248

BIOTECH: 5.0 NEWS HEADLINES: GENOMICS

Main Title: HEALTHCARE REVIEW

Pub. Date: November 01, 2002

Source: DATAMONITOR

Telephone: +44 20 7675 7000

Word Count: 2223 (2 pp.)

Language: English

Country: WORLD

Industry: HEALTH CARE

Company Names (DIALOG Generated): Applera Corporation ; Applied Biosystems Group ; Asterand ; Biologics License Application ; BioMarin Pharmaceutical ; Bristol Myers Squibb ; Celera Diagnostics ; Celera Genomics Group ; Chapel Hill School ; Charles River Laboratories ; Clinical Cancer Research ; Comprehensive Cancer Center ; Genzyme Biosurgery ; Genzyme Corp ; Hybridon ; Isis Pharmaceuticals ; Jackson Laboratory ; Jefferson Medical College ; Leukaemia Research Fund ; Massachusetts General Hospital ; Merck & Co ; National Institute of Allergy and Infectious Diseases ; National Institute of Diabetes Digestive and Kidney Diseases of the National Institutes of Health ; Onyvax ; Pennsylvania School ; Perlegen Sciences ; Pittsburgh Cancer Institute ; Pittsburgh School ; Proteome Systems ; Rockefeller University ; Southern Methodist University ; Stanford University Medical Center ; Target Discovery Program ; Targeted Genetics ; Texas M D Anderson Cancer Center ; Trademark Office ; University of Michigan ; University of North Carolina at Chapel Hill School ; University of Pennsylvania School of Medicine ; University of Pittsburgh Cancer Institute ; University of Pittsburgh School of Medicine ; University of Texas M D Anderson Cancer Center ; UNC ; Wake Forest University School of Medicine ; Wistar Institute

...separate lawsuits, for alleged infringement of its functional genomics US patents.

Hybridon and Aegera form **antisense** drug development collaboration
Hybridon and Aegera Therapeutics have entered into a collaboration and license agreement to research and develop an **antisense** drug targeted to the **XIAP** gene.

Sangamo granted US zinc finger DNA-binding proteins patent
Sangamo BioSciences has been granted...

...in Neuron, could help scientists devise new strategies to block the pain hypersensitivity associated with **inflammation** .

Wake Forest receives \$20 million NIH grant to oversee diabetes genetics study

Wake Forest University...

...illnesses such as cancer and Alzheimer's disease.

Interim results of AVI's phase II **antisense** trial for cardiovascular restenosis confirm safety and efficacy

AVI BioPharma has released positive interim results of a phase II clinical trial measuring the safety and efficacy of its **antisense** compound, Resten-NG, when delivered via catheter during balloon angioplasty procedures.

Xenon Genetics adds two...anorexia nervosa.

Lexicon Genetics discovers new role for protein to develop antibody for treatment of **inflammation**

Lexicon Genetics has discovered a new role in the immune system for a secreted protein that may serve as a target for the development of drugs to treat **inflammation** associated with arthritis and autoimmune disease.

Targeted Genetics cystic fibrosis gene therapy meets primary endpoint...

...key problems in the disease.

Deltagen identifies a novel antibody target for potential treatment of **inflammatory** disorders

Deltagen has discovered a novel drug target, designated DT044I, for the potential treatment of **inflammatory** disorders, including rheumatoid arthritis. This target flows from the company's high-throughput identification screens...

27/K/21 (Item 4 from file: 761)
DIALOG(R) File 761: Datamonitor Market Res.
(c) 2004 Datamonitor. All rts. reserv.

00203191

CANCER: 7.0 NEWS HEADLINES: COLORECTAL

Main Title: HEALTHCARE REVIEW
Pub. Date: November 01, 2002
Source: DATAMONITOR
Telephone: +44 20 7675 7000
Word Count: 3846 (3 pp.)
Language: English

Country: WORLD
Industry: HEALTH CARE
Company Names (DIALOG Generated): Access Pharmaceuticals ; Actinium
Pharmaceuticals ; Adherex Technologies ; Advanced Viral
Research ; Advectus Life Sciences ; American Association for
the Advancement of Science ; Arius Research ; AxCell
Biosciences ; AEterna Laboratories ; Biomira ; Bioniche Life
Sciences ; BioCryst Pharmaceuticals ; Cancer Research ; Cell
Therapeutics ; Chugai Pharmaceutical ; Clinical Cancer Research
; Comprehensive Cancer Center ; Cytogen ; CIMYM ; Dana
Farber/Partners ; FDA 's Office of Orphan Products Development
; Genentech ; Genta ; GenVec ; Guilford Pharmaceuticals ; GPC
Biotech ; Holden Comprehensive Cancer Center ; Hybridon ; HL ;
Idec Pharmaceuticals ; IntraBiotics Pharmaceuticals ; Isis
Pharmaceuticals ; Jackson Laboratory ; King Pharmaceuticals ;
Leland Stanford Junior University ; Leukaemia Research Fund ; M
D Anderson Cancer Center ; Maxim Pharmaceuticals ; Maxygen ;
Mayne Group ; Mayo Clinic Cancer Center ; Medical Devices ;
Memorial Sloan Kettering Cancer Center ; Merck ; Montefiore
Medical Center ; MGI Pharma ; National Cancer Institute ;
NeoTherapeutics ; New Drug Application ; Nippon Shinyaku ;
North Carolina Brain Tumor Center ; Novartis Ophthalmics ;
Orphan Products Development ; Ortho Biotech ; Oxford
GlycoSciences ; OSI Pharmaceuticals ; Peregrine Pharmaceuticals
; Pharmaceutical Research and Manufacturers ; Pittsburgh Cancer
Institute ; Protein Design Labs ; Rockefeller University ;
Royal North Shore Hospital ; Southern Methodist University ;
Stanford University ; SLIL Biomedical ; Texas M D Anderson
Cancer Center ; National Cancer Institute ; Ohio State
University ; University of Iowa ; University of North Carolina
Brain Tumor Center ; University of Texas MD Anderson Cancer
Center ; Trademark Office ; University of California at San
Francisco ; University of North Carolina at Chapel Hill ;
University of Pittsburgh Cancer Institute ; University of Texas
M D Anderson Cancer Center ; University of Wisconsin ;
University of Zurich ; UNC ; Vical ; Vion Pharmaceuticals ;
Viventia Biotech ; Washington University School of Medicine

...a terminal brain cancer for which there is no known cure.
Hybridon and Aegera form **antisense** drug development collaboration
Hybridon and Aegera Therapeutics have entered into a collaboration and
license agreement to research and develop an **antisense** drug targeted to

the **XIAP** gene.

National Institutes of Health funds Seattle Genetics prodrug study
Seattle Genetics has won a...

...and neck radiation therapy used in the treatment of various cancers.

Isis awarded patent for **antisense** compounds

Isis Pharmaceuticals has been issued a US patent covering **antisense** compounds targeted to genes encoding p38 mitogen activated protein kinases or p38 MAP kinases. p38 MAP kinases regulate many biological processes related to **inflammatory** diseases and cancer.

Vical stops Allovectin-7 melanoma program

Vical Incorporated has discontinued its low...

27/K/22 (Item 5 from file: 761)

DIALOG(R) File 761:Datamonitor Market Res.

(c) 2004 Datamonitor. All rts. reserv.

00203178

REVIEW: 36.0 NEWS HEADLINES: ALLIANCES AND JOINT VENTURES

Main Title: GLOBAL HEALTHCARE

Pub. Date: November 01, 2002

Source: DATAMONITOR

Telephone: +44 20 7675 7000

Word Count: 1481 (1 pp.)

Language: English

Country: WORLD

Industry: HEALTH CARE

Company Names (DIALOG Generated): Alza Corporation ; American Heart Association ; Aventis Pasteur ; AAI International ; AM Pharma Holding ; Bayer ; Blue Cross & Blue Shield ; Cambridge Antibody Technology ; Cancer Treatment ; Cel Sci ; Center for Drug Evaluation and Research of the FDA ; Chugai ; Epicyte Pharmaceutical ; Genentech ; Genta ; Hybridon ; Institute 's Molecular Radiation ; Ista Pharmaceuticals ; Kos Pharmaceuticals ; Merck & Co ; National Cancer Institute ; National Institute of Diabetes Digestive and Kidney Diseases of the National Institutes of Health ; Nautilus Biotech ; Naval Medical Research Center of the US Navy ; New Drug Application ; Organogenesis ; Perlegen Sciences ; Pharma ; ProMetic Life Sciences ; Radiation Oncology Sciences Program ; Raptiva ; Senju Pharmaceutical Co ; Speedel Group ; Tanabe Holding America ; Tanabe Seiyaku ; University of North Carolina ; US National Cancer Institute ; Wake Forest University School of Medicine ; Xoma

...treatment of HIV-1 infection in combination with other antiretroviral agents.

Hybridon and Aegera form **antisense** drug development collaboration

Hybridon and Aegera Therapeutics have entered into a collaboration and license agreement to research and develop an **antisense** drug targeted to the **XIAP** gene.

Raptiva achieves primary efficacy endpoint in phase III psoriasis study

Results of a randomized...certain other COX-inhibiting nitric oxide-donators in Japan in the treatment of pain and **inflammation**.

Cambridge Antibody Technology and Chugai enter novel human monoclonal antibodies development partnership
Cambridge Antibody Technology...

27/K/23 (Item 6 from file: 761)

DIALOG(R)File 761:Datamonitor Market Res.
(c) 2004 Datamonitor. All rts. reserv.

00203164

REVIEW: 25.0 NEWS HEADLINES: COLORECTAL

Main Title: GLOBAL HEALTHCARE

Pub. Date: November 01, 2002

Source: DATAMONITOR

Telephone: +44 20 7675 7000

Word Count: 1274 (1 pp.)

Language: English

Country: WORLD

Industry: HEALTH CARE

Company Names (DIALOG Generated): Access Pharmaceuticals ; American Association for the Advancement of Science ; AxCell Biosciences ; Biomira ; Clinical Cancer Research ; Cytogen ; Genta ; GenVec ; Guilford Pharmaceuticals ; Hybridon ; Isis Pharmaceuticals ; Maxim Pharmaceuticals ; Maxygen ; Merck ; National Cancer Institute ; New Drug Application ; Ortho Biotech ; Peregrine Pharmaceuticals ; Pharmaceutical Research and Manufacturers ; Pittsburgh Cancer Institute ; Royal North Shore Hospital ; National Cancer Institute ; University of Pittsburgh Cancer Institute ; Vical ; Vion Pharmaceuticals

...a terminal brain cancer for which there is no known cure.

Hybridon and Aegera form **antisense** drug development collaboration

Hybridon and Aegera Therapeutics have entered into a collaboration and license agreement to research and develop an **antisense** drug targeted to the **XIAP** gene.

National Institutes of Health funds Seattle Genetics prodrug study

Seattle Genetics has won a...

...and neck radiation therapy used in the treatment of various cancers.

Isis awarded patent for **antisense** compounds

Isis Pharmaceuticals has been issued a US patent covering **antisense** compounds targeted to genes encoding p38 mitogen activated protein kinases or p38 MAP kinases. p38 MAP kinases regulate many biological processes related to **inflammatory** diseases and cancer.

Vical stops Allovectin-7 melanoma program

Vical Incorporated has discontinued its low...

27/K/24 (Item 7 from file: 761)

DIALOG(R)File 761:Datamonitor Market Res.
(c) 2004 Datamonitor. All rts. reserv.

00203161

REVIEW: 23.0 NEWS HEADLINES: GENOMICS

Main Title: GLOBAL HEALTHCARE
Pub. Date: November 01, 2002
Source: DATAMONITOR
Telephone: +44 20 7675 7000
Word Count: 1079 (1 pp.)
Language: English

Country: WORLD

Industry: HEALTH CARE

Company Names (DIALOG Generated): Biologics License Application ; BioMarin
Pharmaceutical ; Genzyme Biosurgery ; Genzyme Corp ; Hybridon ;
Isis Pharmaceuticals ; Jackson Laboratory ; Leukaemia Research
Fund ; Massachusetts General Hospital ; Merck & Co ; National
Institute of Allergy and Infectious Diseases ; National
Institute of Diabetes Digestive and Kidney Diseases of the
National Institutes of Health ; Pennsylvania School ; Perlegen
Sciences ; Southern Methodist University ; Stanford University
Medical Center ; University of North Carolina ; University of
Pennsylvania School of Medicine ; UNC ; Wake Forest University
School of Medicine ; Wistar Institute

...separate lawsuits, for alleged infringement of its functional genomics
US patents.

Hybridon and Aegera form **antisense** drug development collaboration
Hybridon and Aegera Therapeutics have entered into a collaboration and
license agreement to research and develop an **antisense** drug targeted to
the **XIAP** gene.

Sangamo granted US zinc finger DNA-binding proteins patent
Sangamo BioSciences has been granted...

...in Neuron, could help scientists devise new strategies to block the pain
hypersensitivity associated with **inflammation** .

Wake Forest receives \$20 million NIH grant to oversee diabetes genetics
study

Wake Forest University...

...illnesses such as cancer and Alzheimer's disease.

Interim results of AVI's phase II **antisense** trial for cardiovascular
restenosis confirm safety and efficacy

AVI BioPharma has released positive interim results of a phase II clinical
trial measuring the safety and efficacy of its **antisense** compound,
Resten-NG, when delivered via catheter during balloon angioplasty
procedures.

Xenon Genetics adds two...

27/K/25 (Item 1 from file: 993)

DIALOG(R)File 993:NewsRoom 2002

(c) 2004 The Dialog Corporation. All rts. reserv.

0513541041 15Y3182J

OTHER NEWS TO NOTE.

BioWorld Today, v13, n179, pNA

Wednesday, September 18, 2002

JOURNAL CODE: AAFH LANGUAGE: English RECORD TYPE: Fulltext

DOCUMENT TYPE: Newsletter

WORD COUNT: 2,759

...validate a number of drug targets associated with cancer. Atugen will develop GeneBlocs, specially designed **antisense** oligonucleotides, which inhibit expression of specific drug target candidates selected by Schering. Though specific financial...

...of Montreal, formed an agreement in which Hybridon will collaborate with Aegea to develop an **antisense** drug candidate targeted to down-regulate Aegea's target, **XIAP**, which has been implicated in the resistance of cancer cells to chemotherapy. The drug candidate...

...Hybridon licensed to Aegea, on a nonexclusive basis, rights to a portfolio of second-generation **antisense** chemistries and oral **antisense** delivery intellectual property owned or licensed by Hybridon. Aegea will pay Hybridon certain collaboration, up...

...a license to certain Isis patents for target validation and functional genomics using first-generation **antisense** oligonucleotides in exchange for undisclosed payments from Sequitur. Subject to a limited right to conclude existing contracts, Sequitur agreed to not practice in the field of second- or next- generation **antisense** oligonucleotides, also known as chimeric **antisense** oligonucleotides.

Keryx Biopharmaceuticals Inc., of Cambridge, Mass., following successful pre-Phase III meetings with the...

...an orally active small- molecule agent that targets a cell-surface protein involved in the **inflammatory** response. The dose-escalation trial will be conducted in up to 30 healthy adult volunteers...

...that Vasogen's immune modulation therapy can significantly reduce cell death induced by lipopolysaccharide, an **inflammatory** stimulus. The attenuation of the **inflammatory** response was associated with an increase in the anti- **inflammatory** cytokine interleukin-10 and a concomitant decrease in the pro- **inflammatory** cytokine interleukin- 1B. The therapy also led to a reduction in the expression of certain enzymes involved in the intracellular response to **inflammation**, including the stress-activated protein kinase c-Jun NH(2)-terminal kinase.

COPYRIGHT 2002 A...

27/K/26 (Item 2 from file: 993)

DIALOG(R) File 993:NewsRoom 2002

(c) 2004 The Dialog Corporation. All rts. reserv.

0505054968 15XL1PPR

Death receptor-mediated apoptosis and the liver

Yoon, Jung-Hwan

Journal of Herpetology, v37, n3, p400

Sunday, September 1, 2002

JOURNAL CODE: AMRR LANGUAGE: English RECORD TYPE: Fulltext

DOCUMENT TYPE: Scholarly Journal ISSN: 0022-1511

WORD COUNT: 7,958

...produced by macrophages including Kupffer cells, monocytes, and T cells in response to infection and **inflammatory** conditions, but also by other cell types, such as B cells, fibroblasts, and hepatocytes. Unlike...

...proteins that regulate the cytochrome c/Apaf-1 caspase activating pathway [23]. Three human IAPs, **XIAP**, c-IAP-1 and c-IAP-2, have been shown to bind procaspase 9 and...

...hepatitis [57]. Fas expression can be induced either by virus-- specific protein expression or by **inflammatory** cytokines such as interleukin-1. Activated CTLs express Fas and induce hepatocyte apoptosis via Fas...
...to eliminate hepatocytes bearing CCC DNA. Since limited apoptosis can delete cells without inducing untoward **inflammatory** reactions, the differential sensitivity of HBV-infected hepatocytes to TRAIL-induced apoptosis could potentially be...

...DRs use caspase 8/10 and Bid to induce apoptosis, targeted caspase inhibition and Bid **antisense** oligonucleotides could be therapeutically useful in cholestatic liver diseases. For example, Bid **antisense** oligonucleotides have already been shown to be therapeutically useful in a rodent model of extrahepatic...strategy to inhibit liver fibrogenesis. Alternatively, apoptosis in the liver by DRs may be pro- **inflammatory**. For example, administration of Fas agonists in the rodent results in liver chemokine expression, which...

...Lugering N, Held J, Domschke W, et al. Caspase activation correlates with the degree of **inflammatory** liver injury in chronic hepatitis C virus infection. Hepatology 2001;34:758-767.

[57] Mochizuki...

...677.

[71] Higuchi H, Miyoshi H, Bronk SF, Zhang H, Dean N, Gores GJ. Bid **antisense** attenuates bile acid-induced apoptosis and cholestatic liver injury. I Pharmacol Exp Ther 2001;299...

...J, Xing Z. Macrophage engulfment of apoptotic neutrophils contributes to the resolution of acute pulmonary **inflammation** in vivo. Am J Respir Cell Mol Biol 1995;12:232-237.

[77] Fadok VA...

...Campe CB, Schrum LW, Rippe RA, et al. Anti-Fas induces hepatic chemokines and promotes **inflammation** by an NF-kappa B-independent, caspase-3-dependent pathway. J Biol Chem 2001;276...

? T S28/MEDIUM,K/ALL

>>>KWIC option is not available in file(s): 398, 399

28/K/1 (Item 1 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

11277558 PMID: 11356694

X-linked inhibitor of apoptosis protein activates the phosphatidylinositol 3-kinase/Akt pathway in rat granulosa cells during follicular development.

Asselin E; Wang Y; Tsang B K

Reproductive Biology Unit and Division of Reproductive Medicine, Department of Obstetrics and Gynecology and Cellular and Molecular Medicine, University of Ottawa, Canada.

Endocrinology (United States) Jun 2001, 142 (6) p2451-7, ISSN 0013-7227 Journal Code: 0375040

Document type: Journal Article
Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

X-linked inhibitor of apoptosis protein (**XIAP**) in granulosa cells is regulated by gonadotropins during follicular development, although the current understanding of the mechanisms by which **XIAP** suppressed granulosa cell apoptosis is incomplete. In the present study, we investigated the possible involvement...

... development and atresia in immature rats, we have demonstrated that gonadotropin treatment increased granulosa cell **XIAP** and phospho-Akt protein contents and suppressed apoptosis. In addition, gonadotropin withdrawal [equine CG (eCG...

... apoptosis and significantly decreased ovarian weight. The increased apoptosis was accompanied by marked decreases in **XIAP** expression and phosphorylation of Akt protein. Infection of granulosa cells from eCG-primed rats with adenoviral sense **XIAP** [lacZ as a control; multiplicity of infection, 1-5] resulted in **XIAP** overexpression and increased phospho-Akt content, whereas **XIAP antisense** expression (multiplicity of infection, 10-40) decreased granulosa cell phospho-Akt level and induced apoptosis...

... the first time the importance and regulation of the PI 3-K survival pathway by **XIAP** in the control granulosa cell apoptosis.

...; Cultured; Chorionic Gonadotropin--administration and dosage--AD; Chorionic Gonadotropin--immunology--IM; Chorionic Gonadotropin--pharmacology--PD; **Follicular Atresia** ; In Situ Nick-End Labeling; Oligodeoxyribonucleotides, **Antisense** --genetics--GE; Ovarian Follicle --metabolism--ME; Phosphorylation; Proteins--genetics--GE; Rats; Rats, Sprague-Dawley; Transfection

Chemical Name: Antibodies; Chorionic Gonadotropin; IAP-like protein, vertebrate; Oligodeoxyribonucleotides, **Antisense** ; Proteins; Proto-Oncogene Proteins; proto-oncogene protein akt; 1-Phosphatidylinositol 3-Kinase

28/K/2 (Item 1 from file: 159)

DIALOG(R) File 159:Cancerlit

(c) format only 2002 Dialog Corporation. All rts. reserv.

02728648 21255627 PMID: 11356694

X-linked inhibitor of apoptosis protein activates the phosphatidylinositol 3-kinase/Akt pathway in rat granulosa cells during follicular development.

Asselin E; Wang Y; Tsang B K

Reproductive Biology Unit and Division of Reproductive Medicine, Department of Obstetrics and Gynecology and Cellular and Molecular Medicine, University of Ottawa, Canada.

Endocrinology (United States) Jun 2001, 142 (6) p2451-7, ISSN 0013-7227 Journal Code: 0375040

Document Type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

X-linked inhibitor of apoptosis protein (**XIAP**) in granulosa cells is

regulated by gonadotropins during follicular development, although the current understanding of the mechanisms by which **XIAP** suppressed granulosa cell apoptosis is incomplete. In the present study, we investigated the possible involvement...

... development and atresia in immature rats, we have demonstrated that gonadotropin treatment increased granulosa cell **XIAP** and phospho-Akt protein contents and suppressed apoptosis. In addition, gonadotropin withdrawal [equine CG (eCG...

... apoptosis and significantly decreased ovarian weight. The increased apoptosis was accompanied by marked decreases in **XIAP** expression and phosphorylation of Akt protein. Infection of granulosa cells from eCG-primed rats with adenoviral sense **XIAP** [lacZ as a control; multiplicity of infection, 1-5] resulted in **XIAP** overexpression and increased phospho-Akt content, whereas **XIAP antisense** expression (multiplicity of infection, 10-40) decreased granulosa cell phospho-Akt level and induced apoptosis...

... the first time the importance and regulation of the PI 3-K survival pathway by **XIAP** in the control granulosa cell apoptosis.

Minor Descriptors: Antibodies--pharmacology--PD; Apoptosis; Cells, Cultured; **Follicular Atresia** ; Gonadotropins, Chorionic--administration and dosage--AD; Gonadotropins, Chorionic--immunology--IM; Gonadotropins, Chorionic--pharmacology--PD; In Situ Nick-End Labeling; Oligodeoxyribonucleotides, **Antisense** --genetics--GE; Ovarian Follicle--metabolism--ME; Phosphorylation; Proteins--genetics--GE; Rats; Rats, Sprague-Dawley; Transfection

Chemical Name: Antibodies; Gonadotropins, Chorionic; IAP-like protein, vertebrate; Oligodeoxyribonucleotides, **Antisense** ; Proteins; Proto-Oncogene Proteins; proto-oncogene protein akt; 1-Phosphatidylinositol 3-Kinase